



## Genotyping protocol

Ect2

IR00004174 / P4174

(ICS internal reference)

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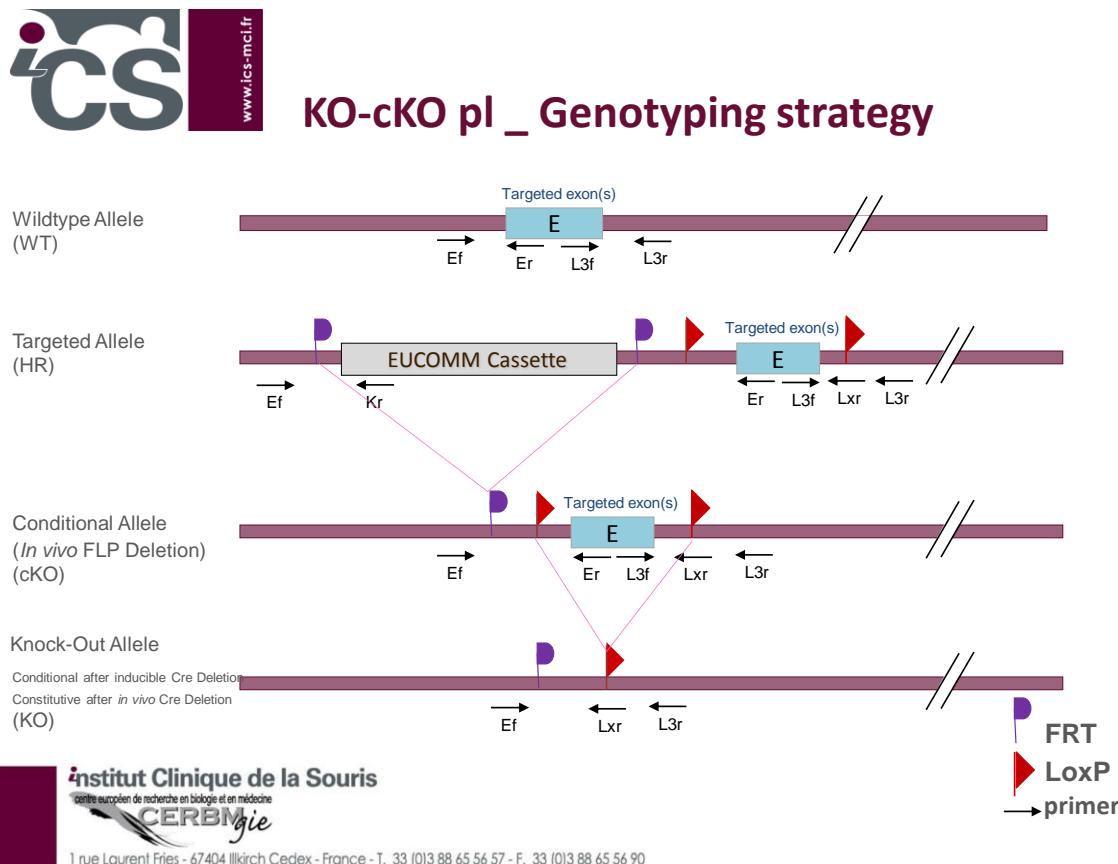
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## 1. Genotyping protocol and data

This section describes the condition used at the Mouse Clinical Institute (ICS) to genotype your **Ect2** Constitutive Knockout / Conditional Knockout (KO-cKO) project.

### 1.1. Genotyping strategy

The map below describes the position of the primers used for genotyping for each possible allele.



## Sequence of primers used for genotyping:

Position	Primers	Sequence
Ef	6724	ATTGTCACTCATGCTAGGGAGGTGC
Er	6727	CGTTGAAGTTCTAGGCTAACCAACAA
Kr	3209	CCAACAGCTTCCCCACAACGG
L3f	6725	TGGCAGGTGGCGAAAGAGCT
L3r	6726	GGCAGCCTGACTGGCACTATTCTTA
Lxr	3255	ACTGATGGCGAGCTCAGACCATAAC

<sup>2</sup>: for a selected position, a second primer was designed

## PCR fragments expected size (bp):

Region analyzed	Primers used	Position on the primer <i>(see the map above)</i>	Targeted allele (HR)	conditional allele (cKO)	WildType allele
5' part of the selection marker	6724-3209	Ef / Kr	318	---	---
Presence of the distal loxP	6725-6726	L3f / L3r	414	414	417
Distal loxP specific PCR	6725-3255	L3f / Lxr	332	332	---
Excision of the selection marker	6724-6727	Ef / Er	5916*	502	402

\*: this PCR product will not be observed using our PCR genotyping conditions (see description below)

\*\*: this PCR is only verified if mice are generated

---: no Amplicon should be obtained

## 1.2. PCR protocol

This section describes the composition of the mix and cycling conditions used for genotyping.

Reagents:

- FastStart PCR Master (Roche)
- DNA (50ng/ $\mu$ l)
- 5' primer (100  $\mu$ M)
- 3' primer (100  $\mu$ M)
- Sterile H<sub>2</sub>O

Volume:

- 7.5 $\mu$ l
- 1.5 $\mu$ l
- 0.06 $\mu$ l
- 0.06 $\mu$ l
- up to 15  $\mu$ l

Cycling conditions:

Temp	Time	#Cycles
95°C	4min	1
94°C	30s	
62°C	30s	34
72°C	1min	
72°C	7min	1
20°C	5min	1

**NB: These PCR conditions have been optimized for high-throughput genotyping. Adaptation to small-scale may be required.**

## 2. Cre and Flp genotyping method

You will find the genotyping protocol in the publication:

[Highly-efficient, fluorescent, locus directed cre and FlpO deleter mice on a pure C57BL/6N genetic background.](#)

Birling MC, Dierich A, Jacquot S, Hérault Y, Pavlovic G.  
Genesis. 2012 Jun;50(6):482-9. doi: 10.1002/dvg.20826. Epub 2012 Mar 20.