



## Genotyping protocol

### Nrp1

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TABLE OF CONTENTS

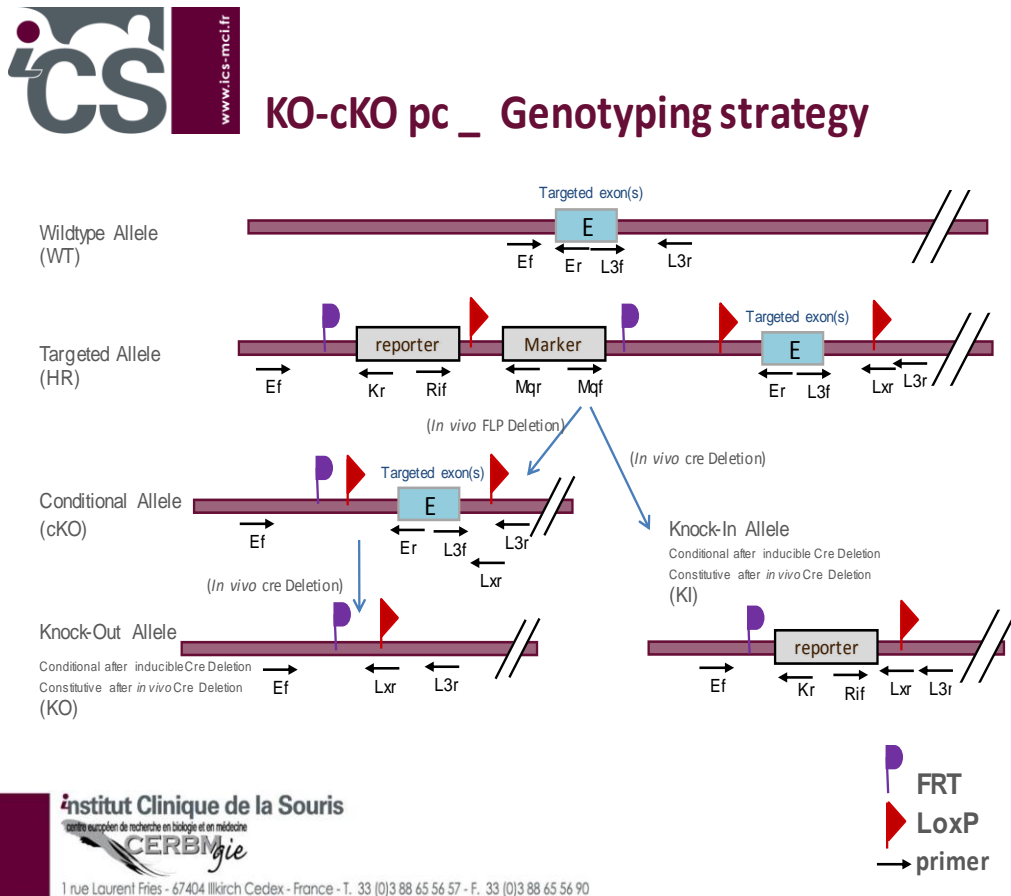
Table of contents .....2  
 1. Genotyping protocol and data .....2  
     1.1. Genotyping strategy .....2  
     1.2. PCR protocol .....4  
 2. Cre and Flp genotyping method .....5

1. Genotyping protocol and data

This section describes the condition used at the Mouse Clinical Institute (ICS) to genotype your Nrp1 Constitutive Knockout / Conditional Knockout (KO-cKO x Cre) 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	8602	GAAAACTAGAGGTCTTGGGGACAGCTG
Er	8605	CCCTAGAAGTTGCTCCCATTTC AAGG
Kr	3278	GGGCAAGAACATAAAAGTGACCCTCC
L3f	8603	CAGAAGTGAAATCCTCATGTCCTGGG
L3r	8604	GATGCAAGAATCACTGGGACTACTGGC
Rif	5966	GCACATGGCTGAATATCGACGGT

### PCR fragments expected size (bp):

Region analyzed	Primers used	Position on the primer (see the map above)	Targeted allele (HR)	conditional allele (cKO)	KI allele	WildType allele
5' part of the selection marker	8602-3278	Ef / Kr	291	---	---	---
Presence of the distal loxP	8603-8604	L3f / L3r	293	293	---	300
Distal loxP specific PCR	8603-3255	L3f / Lxr	151	151	---	---
Excision of the selection marker	8602-8605	Ef / Er	7270*	366	---	243
Cre total excision	5966-3255	Rif / Lxr	3373*	---	471**	---

\*: 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:	Volume:
- FastStart PCR Master (Roche)	7.5µl
- DNA (50ng/µl)	1.5µl
- 5' primer (100 µM)	0.06µl
- 3' primer (100 µM)	0.06µl
- Sterile H <sub>2</sub> O	up to 15 µl

### Cycling conditions:

Temp	Time	#Cycles
95°C	4min	1
94°C	30s	34
62°C	30s	
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.