

# Nphs2-Flox

<b>Nomenclature</b>	C57BL/6Smoc- <i>Nphs2</i> <sup>em1(flox)Smoc</sup>
<b>Cat. NO.</b>	NM-CKO-241788
<b>Strain State</b>	Developing

## Gene Summary

<b>Gene Symbol</b> <b>Nphs2</b>	<b>Synonyms</b>	PDCN; SRN1
	<b>NCBI ID</b>	<a href="#">170484</a>
	<b>MGI ID</b>	<a href="#">2157018</a>
	<b>Ensembl ID</b>	<a href="#">ENSMUSG00000026602</a>
	<b>Human Ortholog</b>	NPHS2

## Model Description

These strains carry loxP sites flanking exon 2 of Nphs2 gene. When crossed with a Cre recombinase-expressing strain, this strain is useful in eliminating tissue-specific conditional expression of Nphs2 gene.

\*Literature published using this strain should indicate: Nphs2-Flox mice (Cat. NO. NM-CKO-241788) were purchased from Shanghai Model Organisms Center, Inc..

## Disease Connection

<b>Nephrotic Syndrome</b>	<b>Phenotype(s)</b>	<a href="#">MGI:4847559</a> Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with CAG-cre mice.
	<b>Reference(s)</b>	Mollet G, Ratelade J, Boyer O, Muda AO, Morisset L, Lavin TA, Kitzi D, Dallman MJ, Bugeon L, Hubner N, Gubler MC, Antignac C, Esquivel EL, Podocin inactivation in mature kidneys causes focal segmental glomerulosclerosis and nephrotic syndrome. J Am Soc Nephrol. 2009 Oct;20(10):2181-9

<b>nephrotic syndrome</b>	<b>Phenotype(s)</b>	<a href="#">MGI:6316681</a> Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with CAG-cre mice.
	<b>Reference(s)</b>	Tabatabaeifar M, Wlodkowski T, Simic I, Denc H, Mollet G, Weber S, Moyers JJ, Bruhl B, Randles MJ, Lennon R, Antignac C, Schaefer F, An inducible mouse model of podocin-mutation-related nephrotic syndrome. PLoS One. 2017;12(10):e0186574

## Validation Data

No data