

# Egln1-Flox

<b>Nomenclature</b>	C57BL/6Smoc- <i>Egln1</i> <sup>em1(flox)Smoc</sup>
<b>Cat. NO.</b>	NM-CKO-2100497
<b>Strain State</b>	Repository Live

## Gene Summary

<b>Gene Symbol</b> Egln1	<b>Synonyms</b>	Phd2; HPH-2; ORF13; SM-20; C1orf12; HIF-PH2; AI503754; Hif-p4h-2
	<b>NCBI ID</b>	<a href="#">112405</a>
	<b>MGI ID</b>	<a href="#">1932286</a>
	<b>Ensembl ID</b>	<a href="#">ENSMUSG00000031987</a>
	<b>Human Ortholog</b>	EGLN1

## Model Description

These mice carry loxP sites flanking exon 2-3 of Egln1 gene. When crossed with a Cre recombinase-expressing strain, this strain is useful in eliminating tissue-specific conditional expression of Egln1 gene.

\*Literature published using this strain should indicate: Egln1-Flox mice (Cat. NO. NM-CKO-2100497) were purchased from Shanghai Model Organisms Center, Inc..

## Disease Connection

<b>Cardiomyopathy</b>	<b>Phenotype(s)</b>	<a href="#">MGI:5304713</a> Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with Egln3-Flox(NM-CKO-2108727) and Myh6-cre mice.
	<b>Reference(s)</b>	Moslehi J, Minamishima YA, Shi J, Neuberg D, Charytan DM, Padera RF, Signoretti S, Liao R, Kaelin WG Jr, Loss of hypoxia-inducible factor prolyl hydroxylase activity in cardiomyocytes phenocopies ischemic cardiomyopathy. Circulation. 2010 Sep 7;122(10):1004-16

## Validation Data

No data

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