

# Fgfr1-Flox

<b>Nomenclature</b>	C57BL/6Smoc- <i>Fgfr1</i> <sup>tm1(flox)Smoc</sup>
<b>Cat. NO.</b>	NM-CKO-240129
<b>Strain State</b>	Developing

## Gene Summary

<b>Gene Symbol</b> Fgfr1	<b>Synonyms</b>	FLG; MFR; Eask; Hspy; Flt-2; c-fgr; FGFR-I; Fgfr-1; AW208770; bFGF-R-1
	<b>NCBI ID</b>	<a href="#">14182</a>
	<b>MGI ID</b>	<a href="#">95522</a>
	<b>Ensembl ID</b>	<a href="#">ENSMUSG00000031565</a>
	<b>Human Ortholog</b>	FGFR1

## Model Description

These strains carry loxP sites flanking exon 8-15 of Fgfr1 gene. When crossed with a Cre recombinase-expressing strain, this strain is useful in eliminating tissue-specific conditional expression of Fgfr1 gene.

\*Literature published using this strain should indicate: Fgfr1-Flox mice (Cat. NO. NM-CKO-240129) were purchased from Shanghai Model Organisms Center, Inc..

## Disease Connection

<b>Atopic Dermatitis</b>	<b>Phenotype(s)</b>	<a href="#">MGI:5642117</a> Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with Fgfr2-Flox(NM-CKO-2101487) and KRT5-cre mice.
	<b>Reference(s)</b>	Sulcova J, Meyer M, Guiducci E, Feyerabend TB, Rodewald HR, Werner S, Mast cells are dispensable in a genetic mouse model of chronic dermatitis. Am J Pathol. 2015 Jun;185(6):1575-87

## Validation Data

No data

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