

Tsc2-Flox

Nomenclature	C57BL/6Smoc- <i>Tsc2</i> ^{tm1(flox)Smoc}
Cat. NO.	NM-CKO-2102032
Strain State	Sperm cryopreservation

Gene Summary

Gene Symbol Tsc2	Synonyms	Tcs2, Nafld
	NCBI ID	22084
	MGI ID	102548
	Ensembl ID	ENSMUSG000000002496
	Human Ortholog	TSC2

Model Description

These mice carry loxP sites flanking exon 3-5 of Tsc2 gene. When crossed with a Cre recombinase-expressing strain, this strain is useful in eliminating tissue-specific conditional expression of Tsc2 gene.

*Literature published using this strain should indicate: Tsc2-Flox mice (Cat. NO. NM-CKO-2102032) were purchased from Shanghai Model Organisms Center, Inc..

Disease Connection

Uterine Fibroid	Phenotype(s)	MGI:5641710 Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with Pgr-Cre mice.
	Reference(s)	Prizant H, Sen A, Light A, Cho SN, DeMayo FJ, Lydon JP, Hammes SR, Uterine-specific loss of Tsc2 leads to myometrial tumors in both the uterus and lungs. Mol Endocrinol. 2013 Sep;27(9):1403-14

Tuberous Sclerosis	Phenotype(s)	MGI:5140838 Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with Pcp2-cre mice.
	Reference(s)	Reith RM, Way S, McKenna J 3rd, Haines K, Gambello MJ, Loss of the tuberous sclerosis complex protein tuberin causes Purkinje cell degeneration. Neurobiol Dis. 2011 Jul;43(1):113-22
tuberous sclerosis	Phenotype(s)	MGI:4880715 Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with GFAP-cre mice.
	Reference(s)	Zeng LH, Rensing NR, Zhang B, Gutmann DH, Gambello MJ, Wong M, Tsc2 gene inactivation causes a more severe epilepsy phenotype than Tsc1 inactivation in a mouse model of Tuberous Sclerosis Complex. Hum Mol Genet. 2011 Feb 1;20(3):445-54

Validation Data

No data