

Wt1-Flox

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| Nomenclature | C57BL/6Smoc- <i>Wt1</i> ^{tm3(flox)Smoc} |
| Cat. NO. | NM-CKO-200041 |
| Strain State | Embryo cryopreservation |

Gene Summary

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|---------------------------|-----------------------|------------------------------------|
| Gene Symbol Wt1 | Synonyms | Wt-1; D630046119Rik |
| | NCBI ID | 22431 |
| | MGI ID | 98968 |
| | Ensembl ID | ENSMUSG00000016458 |
| | Human Ortholog | WT1 |

Model Description

These mice carry loxP sites flanking exon 2-3 of *Wt1* gene. When crossed with a Cre recombinase-expressing strain, this strain is useful in eliminating tissue-specific conditional expression of *Wt1* gene. While *Wt1*-Flox (Stock No. NM-CKO-200234) mice carrying t

Research Application: Cancer research

*Literature published using this strain should indicate: *Wt1*-Flox mice (Cat. NO. NM-CKO-200041) were purchased from Shanghai Model Organisms Center, Inc..

Disease Connection

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| Congenital Diaphragmatic Hernia | Phenotype(s) | MGI:5824267 Note: The expected phenotype(s) may be observed in the above-mentioned mice that bred with <i>Gata4</i> *G2-cre mice. |
| | Reference(s) | Carmona R, Canete A, Cano E, Ariza L, Rojas A, Munoz-Chapuli R, Conditional deletion of WT1 in the septum transversum mesenchyme causes congenital diaphragmatic hernia in mice. <i>Elife</i> . 2016 Sep 19;5:e16009 |

Validation Data

No data
