

# Chd8-KO

|      |                                             |
|------|---------------------------------------------|
| 品系全名 | C57BL/6Smoc- <i>Chd8</i> <sup>em1Smoc</sup> |
| 目录号  | NM-KO-201140                                |
| 品系状态 | 胚胎冻存                                        |

## 基因信息

|             |            |                                                            |
|-------------|------------|------------------------------------------------------------|
| 基因名<br>Chd8 | 基因曾用名      | Chd-8, Duplin, HELSNF1, AU015341, mKIAA1564, 5830451P18Rik |
|             | NCBI ID    | <a href="#">67772</a>                                      |
|             | MGI ID     | <a href="#">1915022</a>                                    |
|             | Ensembl ID | <a href="#">ENSMUSG00000053754</a>                         |
|             | 基因标记细胞类型举例 | 脾脏NK细胞、脂肪组织                                                |

## 品系描述

敲除Chd8基因exon 2-3，建立Chd8基因敲除小鼠模型。

\*使用本品系发表的文献需注明: Chd8-KO mice (Cat. NO. NM-KO-201140) were purchased from Shanghai Model Organisms Center, Inc..

## 疾病预测

|                                     |         |                                                                                                                                                                                                                         |
|-------------------------------------|---------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| 自闭症谱系障碍<br>Autism Spectrum Disorder | 近似模型的表型 | <a href="#">MGI:6189107</a>                                                                                                                                                                                             |
|                                     | 参考文献    | Katayama Y, Nishiyama M, Shoji H, Ohkawa Y, Kawamura A, Sato T, Suyama M, Takumi T, Miyakawa T, Nakayama KI, CHD8 haploinsufficiency results in autistic-like phenotypes in mice. Nature. 2016 Sep 29;537(7622):675-679 |

## 验证数据

暂无数据