

# Ids-KO

|      |                                            |
|------|--------------------------------------------|
| 品系全名 | C57BL/6Smoc- <i>Ids</i> <sup>em1Smoc</sup> |
| 目录号  | NM-KO-200580                               |
| 品系状态 | 活体                                         |

## 基因信息

|                   |            |                                    |
|-------------------|------------|------------------------------------|
| 基因名<br><b>Ids</b> | 基因曾用名      | AW214631                           |
|                   | NCBI ID    | <a href="#">15931</a>              |
|                   | MGI ID     | <a href="#">96417</a>              |
|                   | Ensembl ID | <a href="#">ENSMUSG00000035847</a> |
|                   | 人类同源基因     | IDS                                |
|                   | 人类同源基因关联疾病 | 粘多糖病、肺癌、磺酰脲硫酸酯酶缺乏症                 |

## 品系描述

敲除Ids基因exon 2，建立Ids基因敲除小鼠模型。曾有基因修饰致死报导，详情点击基因信息中的MGI ID。

**应用领域：**黏多糖贮积症(mucopolysaccharidosis, MPS)相关研究

\*使用本品系发表的文献需注明: Ids-KO mice (Cat. NO. NM-KO-200580) were purchased from Shanghai Model Organisms Center, Inc..

## 疾病预测

|                                              |         |                                                                                                                                                                                                                     |
|----------------------------------------------|---------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| 粘多糖贮积症 II<br><b>Mucopolysaccharidosis II</b> | 近似模型的表型 | <a href="#">MGI:3625957</a>                                                                                                                                                                                         |
|                                              | 参考文献    | Cardone M, Polito VA, Pepe S, Mann L, D'Azzo A, Auricchio A, Ballabio A, Cosma MP, Correction of Hunter syndrome in the MPSII mouse model by AAV2/8-mediated gene delivery. Hum Mol Genet. 2006 Apr 1;15(7):1225-36 |

## 验证数据

暂无数据

