B6.129P2(C)-Mecp2<sup>tm1.1bird</sup>/J

Stock No: 003890 | Mecp2<sup>−/−</sup>

 Greenland, Targeted Mutation

**Also Known As: Mecp2<sup>−/−</sup>**

This Mecp2 knockout strain is a mouse model of human Rett Syndrome. Mecp2-deficient males (Mecp2<sup>−/−</sup>) exhibit mobility problems and a range of Rett Syndrome-like characteristics at 3-8 weeks of age. Heterozygous females exhibit a much later onset (~6 months of age).

**Genetic overview**

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<th>N4+N40</th>
<th>(03-APR-18-1)</th>
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Targeted (Null/Knockout) Mecp2 methyl CpG binding protein 2
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