

# Mdr1a/b-Bcrp Knock Out Mouse

**Catalogue number:** 153415

**Sub-type:** Mouse

**Images:**

## Contributor

**Inventor:** Alfred Schinkel

**Institute:** Netherlands Cancer Institute

**Images:**

## Tool details

**\*FOR RESEARCH USE ONLY**

**Name:** Mdr1a/b-Bcrp Knock Out Mouse

**Alternate name:** p-glycoprotein 1, permeability glycoprotein 1, P-gp, pgp, MDR1, ABCB1, CD243 p-glycoprotein 3, BCRP/ABCG2, ATP-binding cassette sub-family G member 2, CDw338,

**Class:**

**Conjugate:**

**Description:** P-gp, a member of the MDR/TAP subfamily, is a glycoprotein encoded in humans by the ABCB1 gene. P-gp is a well-characterized ABC-transporter responsible for transporting a wide variety of substrates across extra- and intracellular membranes. The normal excretion of xenobiotics back into the gut lumen by P-gp pharmacokinetically reduces the efficacy of some pharmaceutical drugs and in addition, some cancer cells also express large amounts of P-gp which can further enhance that effect. This makes some cancers multi-drug resistant.

**Purpose:**

**Parental cell:**

**Organism:**

**Tissue:**

**Model:** Conditional KO

**Gender:**

**Isotype:**

**Reactivity:**

**Selectivity:**

**Host:**

**Immunogen:**

**Immunogen UNIPROT ID:**

**Sequence:**

**Growth properties:**

**Production details:**

This model was generated by breeding the Mdr1a/b mutated mouse with the Bcrp mutated mouse. The Mdr1a/b model was created by sequential targeting of the two Abcb1a and Abcb1b genes in E14 ES cells. Resultant chimeras were backcrossed to FVB/N for seven generations. The Bcrp model was created by targeting the Abcg2 gene in E14 embryonic stem cells derived from 129P2/OlaHsd mice and injecting the targeted cells into FVB blastocysts.

**Formulation:**

**Recommended controls:**

**Bacterial resistance:**

**Selectable markers:**

**Additional notes:** The Mdr1a/b-Bcrp mouse was developed in the laboratory of Alfred Schinkel of the Netherlands Cancer Institute. Useful in studies of drug transport, oral bioavailability and multi-drug resistance

## Target details

**Target:** This model encodes a triple targeted mutation with disruption of the multi-drug resistance genes Abcb1a, Abcb1b and Abcg2.

**Target alternate names:**

**Target background:**

**Molecular weight:**

**Ic50:**

## Applications

**Application:** Useful in studies of drug transport, oral bioavailability and multi-drug resistance

**Application notes:**

## Handling

**Format:**

**Concentration:**

**Passage number:**

**Growth medium:**

**Temperature:**

**Atmosphere:**

**Volume:**

**Storage medium:**

**Storage buffer:**

**Storage conditions:**

**Shipping conditions:** Embryo/Spermatozoa- Dry Ice

## Related tools

**Related tools:** Mrp2 Knock Out Mouse

## References

**References:** van Waterschoot et al. 2010. Br J Pharmacol. 160(5):1224-33. PMID: 20590614. ; Effects of cytochrome P450 3A (CYP3A) and the drug transporters P-glycoprotein (MDR1/ABCB1) and MRP2 (ABCC2) on the pharmacokinetics of lopinavir. ; Vlaming et al. 2006. J Pharmacol Exp Ther. 318(1):319-27. PMID: 16611851. ; Carcinogen and anticancer drug transport by Mrp2 in vivo: studies using Mrp2 (Abcc2) knockout mice.

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