

Genetically engineered models (GEMS)



Mdr1a knockout rat

Model	Mdr1a knockout rat		
Strain	HsdSage: SD -Mdr1a ^{tm1Sage}		
Location	U.S.		
Availability	Live colony		

Characteristics/husbandry

- + Biallelic 20 bp deletion within Abcba1 gene
- + Increased oral bioavailability of P-gp-specific substrates
- + Homozygous knockout rats display total loss of protein via Western blot
- + Background strain: Sprague Dawley

Zygosity genotype

+ Homozygous

Research use

- + DMPK assay
- + PK-PD efflux assay
- + Neurotoxicology
- + Formulation drug-drug interactions
- + Drug resistance
- + Blood brain barrier efflux
- + Efficacy

Origin

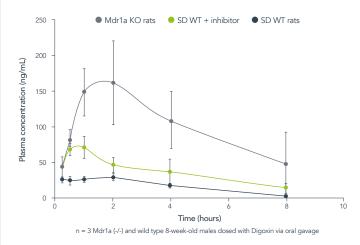
The Mdr1a KO rat model was originally created at SAGE Labs, Inc. in St. Louis, MO and distributed out of the Boyertown, PA facility. The line continues to be maintained through the original SAGE Labs animal inventory acquired by Envigo.

Description

P-glycoprotein plays a critical role in efflux for both brain and liver. Homozygous null Mdr1a rats display increased exposure to CNS drugs in the brain, as well as increased bioavailability in the plasma for P-gp-specific substrates.

MDR1 encodes for P-glycoprotein and is a membrane-bound drug transporter expressed in the brain and intestine. It effectively blocks drugs from crossing the blood-brain barrier. P-gp can confer multiple drug resistance to tumor cells. Absence of P-gp creates a functional deficiency in the blood-brain barrier and results in elevated drug levels in many tissues, making this a useful model for efflux assay, efficacy, formulation, tissue distribution, studying neurotoxicology and chemotherapeutic agents.

Figure 1: Oral absorption of Digoxin in the absence and presence of Quinidine



	Brain	Plasma	Brain:Plasma
Wild type	0	25.5	N.d.
Knockout	339	236	159

*Note: Digoxin concentrations in units of ng/mL

++++

Figure 2: Western blot using proximal colon tissue isolated from both wild type and Mdr1a homozygous knockout tissue

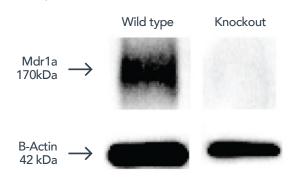
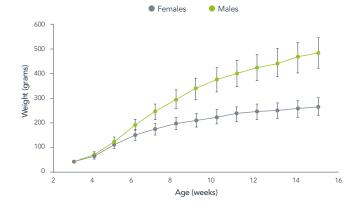


Figure 3: Age/Weight/Curve chart



Citations

29635055 Al-Ali, Ahmed A Abdulhussein; Quach, Jeffrey Rong Chao; Bundgaard, Christoffer; Steffansen, Bente; Holm, René; Nielsen, Carsten Uhd; Polysorbate 20 alters the oral bioavailability of etoposide in wild type and mdr1a deficient Sprague-Dawley rats. International journal of pharmaceutics Vol.543, 2018

22947335 Dulin, Jennifer N; Moore, Meredith L; Grill, Raymond J; The dual cyclooxygenase/5-lipoxygenase inhibitor licofelone attenuates p-glycoprotein-mediated drug resistance in the injured spinal cord. Journal of neurotrauma Vol.30, 2013

26105563 Dumas, Noé; Moulin-Sallanon, Marcelle; Fender, Pascal; Tournier, Benjamin B; Ginovart, Nathalie; Charnay, Yves; Millet, Philippe; In Vivo Quantification of 5-HT2A Brain Receptors in Mdr1a KO Rats with 123I-R91150 Single-Photon Emission Computed Tomography. Molecular imaging Vol.14, 2015

25053619 Fuchs, Holger; Kishimoto, Wataru; Gansser, Dietmar; Tanswell, Paul; Ishiguro, Naoki; Brain penetration of WEB 2086 (Apafant) and dantrolene in Mdr1a (P-glycoprotein) and Bcrp knockout rats. Drug metabolism and disposition: the biological fate of chemicals Vol.42, 2014 26431714 Fujii, Shinobu; Setoguchi, Chikako; Kawazu, Kouichi; Hosoya, Ken-ichi; Functional Characterization of Carrier-Mediated Transport of Pravastatin across the Blood-Retinal Barrier in Rats. Drug metabolism and disposition: the biological fate of chemicals Vol.43, 2015

24985475 Fujii, Shinobu; Setoguchi, Chikako; Kawazu, Kouichi; Hosoya, Ken-ichi; Impact of P-glycoprotein on blood-retinal barrier permeability: comparison of bloodaqueous humor and blood-brain barrier using mdr1a knockout rats. Investigative ophthalmology & visual science Vol.55, 2014

29674491 Ganguly, Samit; Panetta, John C; Roberts, Jessica K; Schuetz, Erin G; Ketamine Pharmacokinetics and Pharmacodynamics Are Altered by P-Glycoprotein and Breast Cancer Resistance Protein Efflux Transporters in Mice. Drug metabolism and disposition: the biological fate of chemicals Vol.46, 2018

25539457 Huang, Liyue; Li, Xingwen; Roberts, Jonathan; Janosky, Brett; Lin, Min-Hwa Jasmine; Differential role of P-glycoprotein and breast cancer resistance protein in drug distribution into brain, CSF and peripheral nerve tissues in rats. Xenobiotica; the fate of foreign compounds in biological systems Vol.45, 2015

24398459 Liu, Xingrong; Cheong, Jonathan; Ding, Xiao; Deshmukh, Gauri; Use of cassette dosing approach to examine the effects of P-glycoprotein on the brain and cerebrospinal fluid concentrations in wild-type and P-glycoprotein knockout rats. Drug metabolism and disposition: the biological fate of chemicals Vol.42, 2014

31685755 Miyake, Taiji; Estimating Efflux Transporter-Mediated Disposition of Molecules beyond the Rule of Five (bRo5) Using Transporter Gene Knockout Rats. Biological & pharmaceutical bulletin Vol.43, 2020

26830080 Nagaya, Yoko; Nozaki, Yoshitane; Takenaka, Osamu; Watari, Ryuji; Kusano, Kazutomi; Yoshimura, Tsutomu; Kusuhara, Hiroyuki; Investigation of utility of cerebrospinal fluid drug concentration as a surrogate for interstitial fluid concentration using microdialysis coupled with cisternal cerebrospinal fluid sampling in wild-type and Mdr1a(-/-) rats. Drug metabolism and pharmacokinetics Vol.31, 2016

31678424 Nicolas, Jean-Marie; Chanteux, Hugues; Nicolaï, Johan; Brouta, Frédéric; Viot, Delphine; Rosseels, Marie-Luce; Gillent, Eric; Bonnaillie, Pierre; Mathy, François-Xavier; Long, Jeff; Helmer, Eric; Role of P-glycoprotein in the brain disposition of seletalisib: Evaluation of the potential for drug-drug interactions. European journal of pharmaceutical sciences : official journal of the European Federation for Pharmaceutical Sciences Vol.142, 2020

27601334 Nielsen, Carsten Uhd; Abdulhussein, Ahmed A; Colak, Dilan; Holm, René; Polysorbate 20 increases oral absorption of digoxin in wild-type Sprague Dawley rats, but not in mdr1a(-/-) Sprague Dawley rats. International journal of pharmaceutics Vol.513, 2016

31682973 Sato, Sho; Tohyama, Kimio; Kosugi, Yohei; Investigation of MDR1overexpressing cell lines to derive a quantitative prediction approach for brain disposition using in vitro efflux activities. European journal of pharmaceutical sciences : official journal of the European Federation for Pharmaceutical Sciences Vol.142, 2020

24839994 Suzuki, Motoya; Komura, Hiroshi; Yoshikawa, Tomonori; Enya, Seiji; Nagao, Akemi; Takubo, Hiroaki; Kogayu, Motohiro; Characterization of gastrointestinal absorption of digoxin involving influx and efflux transporter in rats: application of mdr1a knockout (-/-) rats into absorption study of multiple transporter substrate. Xenobiotica; the fate of foreign compounds in biological systems Vol.44, 2014

22711747 Zamek-Gliszczynski, Maciej J; Bedwell, David W; Bao, Jing Q; Higgins, J William; Characterization of SAGE Mdr1a (P-gp), Bcrp, and Mrp2 knockout rats using loperamide, paclitaxel, sulfasalazine, and carboxydichlorofluorescein pharmacokinetics. Drug metabolism and disposition: the biological fate of chemicals Vol.40, 2012

23569176 Zamek-Gliszczynski, Maciej J; Goldstein, Keith M; Paulman, April; Baker, Thomas K; Ryan, Timothy P; Minor compensatory changes in SAGE Mdr1a (P-gp), Bcrp, and Mrp2 knockout rats do not detract from their utility in the study of transportermediated pharmacokinetics. Drug metabolism and disposition: the biological fate of chemicals Vol.41, 2013

Contact us



North America 800.793.7287 gemsorders@envigo.com Envigo, 8520 Allison Pointe Blvd., Suite 400, Indianapolis, IN 46250, United States

© 2020 Envigo.