



Oncology Animal Models

The following information provides an overview of Charles River's portfolio of immunodeficient and immunocompetent oncology animal models produced in North America.

Immunodeficient Animal Models

Immunodeficient animal models are extremely useful in a wide range of biomedical research, including infectious disease, stem cell, immunology and oncology studies. Due to the unique vulnerability that makes these models vital to research, their care and maintenance demands a high level of expertise and technological resources.

| Strain | Hair | T-Cell Deficient | B-Cell Deficient | NK Cell Deficient |
|-------------------------------------|------|------------------|------------------|-------------------|
| Athymic Nude Mouse | No | Yes | No | No |
| CD-1® Nude Mouse | No | Yes | No | No |
| NU/NU Mouse | No | Yes | No | No |
| BALB/c Nude Mouse | No | Yes | No | No |
| NIH-III Mouse | No | Yes | Yes | Impaired |
| RNU Rat | No | Yes | No | No |
| SCID Hairless Outbred (SHO®) Mouse | No | Yes | Yes | No |
| SCID Hairless Congenic (SHC™) Mouse | No | Yes | Yes | No |
| Fox Chase SCID® Congenic Mouse | Yes | Yes | Yes | No |
| Fox Chase SCID® Beige Mouse | Yes | Yes | Yes | Impaired |
| NOD SCID Mouse | Yes | Yes | Yes | Impaired |

Nude Models

Athymic Nude Mouse

Nomenclature: Crl:NU(NCr)-*Foxn1^{nu}* **Strain Code:** 490 (homozygous), 491 (heterozygous) **Origin:** This immunodeficient nude mouse originated from NIH and was originally thought to be a BALB/c congenic. It was later determined that it was not inbred and is, therefore, maintained as an outbred. This model is not associated with any stock or strain. The animal lacks a thymus, is unable to produce T cells, and is therefore immunodeficient. To Charles River from NCI in 2010.

Overview

Oncology is one of the leading areas of research into new therapeutics. Due to the challenges inherent in researching and developing anticancer therapeutics, it is critical that you have the right tools and resources available to you. Backed by decades of technical, scientific and veterinary experience, Charles River's global portfolio of high-quality oncology models gives you the benefit of partnering with an industry leader offering an infrastructure capable of advancing your research now and in the future.



CD-1® Nude Mouse

Nomenclature: Crl:CD1-*Foxn1^{nu}* **Strain Code:** 086 (homozygous), 087 (heterozygous) **Origin:** Developed from the transfer of the nude gene from Crl:NU-*Foxn1^{nu}* to a CD-1 mouse through a series of crosses and backcrosses beginning in 1979 at Charles River Wilmington, MA. The animal lacks a thymus, is unable to produce T cells and is therefore immunodeficient.

NU/NU Mouse

Nomenclature: Crl:NU-*Foxn1^{nu}* **Strain Code:** 088 (homozygous), 089 (heterozygous) **Origin:** This immunodeficient nude mouse originated from NIH and was originally thought to be a BALB/c congenic. It was later determined that it was not inbred and is, therefore, maintained as an outbred. This model is not associated with any stock or strain. The animal lacks a thymus, is unable to produce T cells and is therefore immunodeficient.

BALB/c Nude Mouse

Nomenclature: CAnN.Cg-*Foxn1^{nu}*/Crl **Strain Code:** 194 (homozygous), 195 (heterozygous) **Origin:** Developed through crosses and backcrosses between BALB/cABom-nu and BALB/cAnNCrj-nu at Charles River Japan. Pedigreed pregnant females of CAnN.Cg-*Foxn1^{nu}*/Crl were received from Charles River Japan in 1985. This mouse is inbred, and genetic monitoring results confirm it to be a BALB/c nude. The homozygous animal lacks a thymus, is unable to produce T cells and is therefore immunodeficient.

NIH-III Mouse

Nomenclature: Crl:NIH-*Lyst^{tg}Foxn1^{nu}Btk^{xid}* **Strain Code:** 201 (homozygous), 202 (heterozygous) **Origin:** Most commonly called the NIH-III, it was developed at NIH. In addition to the nude gene, which results in the absence of thymus and T-cell function, this mouse has two other mutations important in regulating the function of the immune system. These are designated as x-linked immune defect (*xid*) and beige (*bg*). The *xid* mutation affects the maturation of T-independent B lymphocytes. It has been demonstrated that *bg* homozygotes have defective natural killer (NK) cells that are cytotoxic *in vitro* to tumor cells. However, the extent of the T-independent B-lymphocyte and NK cell deficiencies in the NIH-III have not been established.

RNU Nude Rat

Nomenclature: Crl:NIH-*Foxn1^{nu}* **Strain Code:** 316 (homozygous), 118 (heterozygous) **Origin:** The NIH nude rat was developed in 1979–1980 through a series of matings involving eight inbred rat strains. To Charles River from the National Institutes of Health in 2001. This athymic nude rat is T-cell deficient and shows depleted cell populations in thymus-dependent areas of peripheral lymphoid organs.

Severe Combined Immunodeficiency (SCID) Models

SCID Hairless Outbred (SHO®) Mouse

Nomenclature: Crl:SHO-*Prkdc^{scid}Hr^{hr}* **Strain Code:** 474 **Origin:** The hairless SCID mouse was produced by Charles River Research Models in 2007 by intercrossing the Crl:HA-*Prkdc^{scid}* and Crl:SKH1-*Hr^{hr}* stocks. The resulting animals are homozygous for the *Prkdc^{scid}* and the *Hr^{hr}* mutations and thus exhibit the severe combined immunodeficiency phenotype characteristic of SCID mice and are also hairless.

SCID Hairless Congenic (SHC™) Mouse

Nomenclature: CB17.Cg-*Prkdc^{scid}Hr^{hr}*/lcrCrl **Strain Code:** 488 **Origin:** The hairless SCID congenic was created in 2009 by Charles River Research Models by using marker-assisted accelerated backcrossing to place the hairless gene (*Hr^{hr}*) present in the Crl:SKH1-*Hr^{hr}* stock onto a CB17/lcr-*Prkdc^{scid}*/lcrCrl genetic background. These mice are homozygous for both *Hr^{hr}* and *Prkdc^{scid}* mutations, so exhibit the severe combined immunodeficiency phenotype characteristic of SCID mice and are also hairless.



Fox Chase SCID® Congenic Mouse

Nomenclature: CB17/lcr-*Prkdc^{scid}*/lcrIcoCrl **Strain Code:** 236 **Origin:** SCID mice possess a genetic autosomal recessive mutation (SCID). Discovered in 1980 by Bosma in C.B-17/lcr mice at Fox Chase Cancer Center. SCID mice show a severe combined immunodeficiency affecting both B and T lymphocytes. They have normal natural killer (NK) cells, macrophages, and granulocytes. To Charles River in 1991 from a Charles River France foundation colony.



Fox Chase SCID® Beige Mouse

Nomenclature: CB17.Cg-*Prkdc^{scid}**Lyst^{bg}*/Crl **Strain Code:** 250 **Origin:** A congenic mouse that possesses both autosomal recessive mutations SCID (*Prkdc^{scid}*) and beige (*Lyst^{bg}*). The SCID mutation results in severe combined immunodeficiency affecting both the B and T lymphocytes. The beige mutation results in defective natural killer (NK) cells. This mouse was developed by Croy, *et al.*, at the University of Guelph by an intercross of C.B-17 *scid/scid* to C57BL/6 *bg/bg* mice. To Charles River in 1993.



NOD SCID Mouse

Nomenclature: NOD.CB17-*Prkdc^{scid}*/NcrCrl **Strain Code:** 394 **Origin:** The SCID mutation has been transferred onto a non-obese diabetic background. Animals homozygous for the SCID mutation have impaired T- and B-cell lymphocyte development and impaired natural killer (NK) cells. To Charles River in 2005 from Frederick Cancer Research and Development Center.

Immunocompetent Models

Buffalo Rat

Nomenclature: BUF/CrCrl **Strain Code:** 281 **Origin:** Heston in 1946 from Buffalo stock of H. Morris. To NIH in 1951 at F10. To Charles River in 1998 from the National Cancer Institute Animal Production Program (Cr). **Coat Color:** White (Albino).

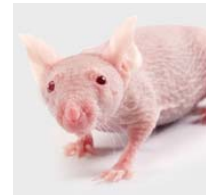
Copenhagen Rat

Nomenclature: COP/CrCrl **Strain Code:** 286 **Origin:** Curtis in 1921 at Columbia University Institute for Cancer Research. To National Cancer Institute Animal Production Program (Cr). To Charles River from the National Cancer Institute in 1998. **Coat Color:** White with a brown hood.

p53 TGEM® Knockout Rat

Nomenclature: CrI:WI(UL)-*Tp53^{tm1/Hubr}* **Origin:** At the Hubrecht Institute, a nonsense mutation at amino acid position 273 (Cys to stop) within the DNA binding domain of the rat p53 gene was isolated, which resulted in a full knockout mutation. Systematic generation of the TGEM® knockout rats was carried out by random mutagenesis of Wistar rats followed by PCR amplification and capillary sequencing. Transferred to Charles River under exclusive license from Transposagen in 2010. **Coat Color:** White (Albino).





Immortomouse®*

Nomenclature: CBA;B10-Tg(H2K^b-tsA58)6Kio/CrI **Strain Code:** 237 (homozygous), 238 (hemizygous) **Origin:** At the Ludwig Institute for Cancer Research, a hybrid construct containing H-2K^b (MHC Class I antigen) 5' promotor sequences fused to the early region of the SV40 mutant tsA58, which encodes both the large and small SV40 tumor antigens, was microinjected into fertilized oocytes from CBA/Ca x C57BL/10 F1 mice. Following reimplantation, 88 mice were born, of which 34 were transgenic and carried one to five copies of the gene. RNA from a variety of tissues from one nontransgenic and three transgenic animals was analyzed by Northern blot analysis using a SV40 early region-specific probe. RNA extracted from tissues of transgenic mice contained varying amounts of a 2.5kb RNA species, while no tsA58 TAG RNA was detected in tissues of the nontransgenic mouse; thymus and liver showed the highest level of expression, while brain showed the lowest. Distribution rights to Charles River in 1991. **Coat Color:** Primarily agouti, infrequently black.

*Transgenic, isolator-maintained

Cryopreserved Oncology Models

The strains listed below are currently maintained as frozen embryos. Two breeding pairs can be made available upon request approximately 12 weeks after order receipt. A dedicated supply can also be established for larger order requests. Please contact Customer Service for pricing and availability.

| Common Name | Nomenclature | Species | Coat Color |
|----------------|--|---------|----------------------------|
| AKR | AKR/NCrI | Mouse | White (Albino) |
| BDIX | BDIX/CrCrI | Rat | Agouti |
| Noble | NBL/CrCrI | Rat | White with black face/hood |
| Wistar Furth | WF/CrCrI | Rat | Albino |
| B6 D933A | B6J.129P2-Pik3ca ^{tm2Bvan} /CrI | Mouse | Black |
| p110alpha flox | B6J.129P2-Pik3ca ^{tm1Bvan} /CrI | Mouse | Black |
| B6 LacZ | B6J.129P2-Pik3ca ^{tm3Bvan} /CrI | Mouse | Black |
| p110beta flox | B6J.129P2-Pik3cb ^{tm1Bvan} /CrI | Mouse | Black |
| B6 lacZ D931A | B6J.129P2-Pik3cb ^{tm2Bvan} /CrI | Mouse | Black |
| B6 D931A | B6J.129P2-Pik3cb ^{tm3Bvan} /CrI | Mouse | Black |
| 129 D931A | 129P2.B6J-Pik3cb ^{tm3Bvan} /CrI | Mouse | Agouti |
| C D931A | C.129P2(B6)-Pik3cb ^{tm3Bvan} /CrI | Mouse | White (Albino) |
| B6 flox | B6J.129P2-Pik3cb ^{tm1.1Bvan} /CrI | Mouse | Black |
| 129 flox | 129P2;B6J-Pik3cb ^{tm1.1Bvan} /CrI | Mouse | Agouti |
| lacZ | B6J.129P2-Pik3cd ^{tm1Bvan} /CrI | Mouse | Black |
| B6D910A | B6J.129P2-Pik3cd ^{tm2Bvan} /CrI | Mouse | Black |
| C D910A | C.129P2(B6)-Pik3cd ^{tm2Bvan} /CrI | Mouse | Agouti |
| B6 S1031A | B6J.129P2-Pik3cd ^{tm3Bvan} /CrI | Mouse | Black |