



Hybrid Mice

B6C3F1 (C57BL/6 x C3H)F1

Origin

B6C3F1/OlaHsd

This F1 hybrid is a cross between C57BL/6JOlaHsd females and C3H/HeNHsd males.

B6C3F1/Hsd

This F1 hybrid is a cross between C57BL/6NHsd females and C3H/HeNHsd males.

Research application

Toxicology, carcinogenic studies, embryo donors in transgenic research.

Characteristics

The F1 hybrid of two inbred strains can be a useful animal for many purposes. It is genetically uniform and heterozygous for all the genes for which the two parental strains differ. F1 animals are easy to produce (hybrid vigour) and are less susceptible to environmental influences than the parent inbred strain. F1 mice will accept transplants of tissues from mice of either parental strain.

Carcinogens

The neonatal B6C3F1 mouse tumorigenicity bioassay is highly sensitive to direct-acting genotoxic carcinogens, with the liver being the principal target organ. (Flammang *et al*, 1997; Fu *et al*, 1997; Fu *et al*, 1998).

Genetics

Coat color genes - a/A, B/B, C/C, D/D : agouti.

Histocompatibility - H-2^{b/k}.

The B6C3F1 will be heterozygous for all the loci where the C57BL/6 and C3H differ and homozygous for all the loci where both parental strains are the same.

Life-span and spontaneous disease

Neoplastic and nonneoplastic lesions in ageing B6C3F1 mice have been described by Ward *et al* (1979). Urinary calculi in the bladder of a male mouse have been described by Wojcinski *et al* (1992). The mean life-span of offspring from reciprocal crosses between C3H/HeJ mice with Mammary Tumor Virus and C57BL/6J have been described by Storer (1966).

Mean life-span of C3B6F1 females is 13,8 months and of C3B6F1 males is 26.3 months, whereas the mean life-span of B6C3F1 females is 27.8 months and of B6C3F1 males is 30.9 months (Storer, 1966).

Mean life-span of B6C3F1 females is 29.9 months and of B6C3F1 males is 30.9 months (Meyers, 1978). Survival and growth patterns have been described by Cameron *et al* (1985). Spontaneous neoplasms have been described by Chandra and Frith (1992). A cellular oncogene was found in spontaneous liver tumors (Fox and Watanabe, 1985). Spontaneous vascular endothelial cell tumors in aged B6C3F1 mice have been described by Yamate *et al* (1988). Evaluating body weight markers for individual animals, as opposed to mean values in an experiment of 50 animals has been shown to be effective in predicting the risk of certain chronic diseases (Seilkop, 1995).

Miscellaneous

Effects of restraint, cage transportation, anesthesia and repeated bleeding on plasma glucose levels have been described by Tabata *et al* (1998).

Nutrition

Caloric restriction and resistance to environmental disease have been described by Frame *et al* (1998).

Physiology and biochemistry

Exposure to electromagnetic fields of 902 MHz and 1747 MHz does not cause a measurable increase in rectal body temperature or in the corticosterone level of B6C3F1 mice secured in fixation tubes (Kamlage, 2002).

References

- Cameron TP, Hickman RL, Kornreich MR, Tarone RE (1985) History, survival, and growth patterns of B6C3F1 mice and F344 rats in the National Cancer Institute carcinogenesis testing program. Fundam. Appl. Toxicol. 5, 526-538.
- Chandra M, Frith CH (1992) Spontaneous neoplasms in B6C3F1 mice. Toxicology Letters 6, 91.
- Fox TR, Watanabe PG (1985) Detection of a cellular oncogene in spontaneous liver tumors in B6C3F1 mice. Science 228, 596-597.
- Flammang TJ, Von Tungeln LS, Kadlubar FF, Fu PP (1997) Neonatal mouse bioassay for tumorigenicity: Alternative to the chronic rodent bioassay. RegulatoryToxicology and Pharmacology. 26, 230-240.
- Frame LT, Hart RW, Leaky JEA (1998) Caloric restriction as a mechanism mediating resistance to environmental disease. Environ. Health Perspect 106, 313-324.
- Fu PP, Von Tungeln LS, Hammons GJ, McMahon G, Wogan GN, Flammang, Kadlubar FF (1997) Carcinogenicity in the B6C3F1 neonatal mouse: a potential alternative bioassay. In: Progress in Clinical and Biological Research. (LaBoef RA, Slaga TJ. Tennant R, eds). Wiley-Liss series.

- Fu PP, Von Tungeln LS, Ping Y, Xia Q, Casciano D, Flammang, Kadlubar FF (1998) Neonatal mouse tumorgenicity bioassay. Drug Information Journal 32, 711-728.
- Kamlage M (2002) Beeinflussung der Körpertemperatur und des vasalen Kortikosteronspiegels durch hochfrequente elektromagnetische Felder des Mobilfunks (902 MHz und 1747 MHz) bei B6C3F1-Mäusen. Dissertation, Hannover, Tierärztliche Hochschule.
- Myers DD (1978) Review of disease patterns and life span in aging mice: Genetic and environmental interactions. Birth defects: Original article series 14, 43-51.
- Seilkop SK (1995) The effect of body weight on tumor incidence and carcinogenicity testing in B6C3F1 mice and F344 rats. Fundam. Appl. Toxicol. 24, 247-259.
- Storer JB (1966b) Nonspecific life shortening in male mice exposed to the mammary tumor agent. J. Natl. Cancer Inst. 37, 211-215.
- Tabata H, Kitamura T, Nagamatsu N (1998) Comparison of restraint, cage transportation, anesthesia and repeated bleeding on plasma glucose levels between mice and rats. Lab. Anim. 32, 142-148.

- Turturro A, Duffy PH Hart RW, Allaben WT (1996) Rational use of dietary control in toxicity studies – B6C3F1 mouse. Toxicol. Pathol. 24, 769-775.
- Ward JM, Goodman DG, Squire RA, Chu KC, Linhart MS (1979) Neoplastic and nonneoplastic lesions in aging (C57BL/6N x C3H/HeN)F1 (86C3F1) mice. J. Natl. Cancer Inst. 63, 849-853.
- Wojcinski ZW, Renlund RC, Barsoum NJ, Smith GS (1992) Struvite urolithiasis in a B6C3F1 mouse. Lab. Anim. 26, 281-287.
- Yamate J, Tajima M, Ihara M, Shibuya K, Kudow S (1988) Spontaneous vascular endothelial cell tumors in aged B6C3F1 mice. Jap. J. Vet. Sci. 50, 453-461.

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