

Research and Reviews: Journal of Zoological Sciences

Laboratory Mice in Medical Research

Babita Seliya*

Department of Biotechnology, Graphic Era University, Dehradun, India

Review Article

Received: 23/08 /2016

Revised: 29/08 /2016

Accepted: 23/09 /2016

*For Correspondence

Department of Biotechnology, Graphic Era University, Dehradun, India, Tel: +919997095117

E-mail: babitaseliya@gmail.com

Keywords: Immunodeficient nude mice, C57BL/6, Transgenic mice.

ABSTRACT

The science of mice has been examined for quite a long time with the fundamental motivation behind irritation control. With the coming of exploratory ways to deal with human medicinal consideration, mice have turned into the most famous creature model framework as a result of the numerous likenesses between their physiology and that of people. All the more as of late, they have been utilized as a part of a considerably more extensive scope of investigative studies went for controlling their genome. With the advancement of olfactory neuroscience from 1990s and sequencing the entire mouse genome toward the start of 21st century, olfactory correspondence is currently learned at cell and atomic levels. The information of mouse science helps in ideally outlining mouse behavioural trials additionally mouse lodging conditions and keeps away from pollution of olfactory flagging variables that may influence the outcomes fundamentally. These laboratory mice have lots of application in Medical Research as well as in novel drug discovery.

INTRODUCTION

Mice assume a significant part in biomedical exploration. Around 95 per cent of all lab creatures are mice and rats. Decreasing dependence on higher-request species, mice have turned into the creature model of decision for biomedical specialists in light of the fact that their physiology ^[1] and hereditary ^[2] make-up nearly takes after that of individuals. Regardless of specific contrasts amongst individuals and rodents, the likenesses are sufficiently solid to give specialists a gigantically capable and adaptable mammalian framework in which to explore human malady.

The sequencing of rat genomes has empowered analysts to reproduce human infections ^[3, 4] in mice through hereditary designing. Scientists are now capable in recognising the illness related attributes in mice and rats, and new innovation permits specialists to specifically alter the DNA of the mice. Research with hereditarily adjusted mice and rats has prompted noteworthy new medications, cures and treatments and keeps on reforming science and pharmaceutical ^[5, 6].

HISTORY

Mice had been used in biomedical study on account that the 16th Century when William Harvey used them for his experiences on blood circulation and reproduction Robert Hooke used them to investigate the organic consequences of an increase in air strains throughout the 18th Century Joseph Priestley and Antoine Lavoisier both used mice to learn breathing. Within the 19th Century Gregor Mendel ^[7] carried out his early investigations of inheritance on mouse coat colour however used to be asked with the aid of his superior to discontinue breeding in his telephone "stinky creatures that, moreover, copulated and had sex". He then switched his investigations to peas but, as his observations have been published in a moderately vague botanical journal, they have been essentially left out for over 35 years except they have been rediscovered in the early 20th Century. In 1902 Lucien Cuénow not published the results of his experiments using mice which showed that Mendel's legal guidelines of inheritance have been also legitimate for animal's results ^[8-9] that have been quickly proven and accelerated to different species.

In the early part of the 20th century Clarence prepare dinner Little, a Harvard undergraduate used to be conducting reviews on mouse genetics ^[10] within the laboratory of William Ernest castle. Little and citadel collaborated intently with Abbie Lathrop who was once a breeder of fancy mice and rats which she marketed to rodent hobbyists and keepers of uncommon pets, and later began selling in colossal numbers to scientific researchers. Together they generated the DBA (Dilute, Brown and non-Agouti) inbred mouse pressure and initiated the systematic generation ^[11, 12] of inbred traces. The mouse has considering the fact that been used generally as a mannequin organism and is associated with many essential biological discoveries of the 20th and 21st Centuries.

Genome

Sequencing of the laboratory mouse genome used to be accomplished in late 2002 making use of the C57BL/6 strain. This used to be only the second mammalian genome ^[13-15] to be sequenced after humans. The haploid genome is set three billion base pairs long (3,000 Mb distributed over 20 chromosomes), for this reason equal to the size of the human genome ^[16-19]. Estimating the number of genes contained in the mouse genome is difficult, partly for the reason that the definition of a gene is still being debated and elevated. The current count of main coding genes within the laboratory mouse is 23,139. Compared to an estimated 20,774 in humans.

Transgenic Strains and Mutant

More than a few mutant lines ^[20] of mice were created by using a quantity of approaches. A small selection from the numerous available strains entails -

Mice attributable to ordinary breeding:

- Non-obese diabetic (NOD) mice, which increase diabetes mellitus, style 1.
- Murphy Roths colossal (MRL) mice with exotic regenerative capacities
- "Waltzing" mice, which stroll in a circular pattern as a result of a mutation adversely affecting their inner ears
- Immunodeficient nude mice, missing thymus and a hair: The mice don't produce T lymphocytes; accordingly, don't mount cellular immune responses. They are used for research in transplantation and immunology ^[21].
- extreme mixed immunodeficient, with an almost completely faulty immune approach

Transgenic mice ^[22, 23], with overseas genes inserted into their genome:

- Abnormally large mice, with an inserted rat development hormone gene
- Onco mice, with an activated oncogene, so to enormously increase the incidence of cancer
- Doogie mice, with improved NMDA receptor function, leading to accelerated reminiscence and learning ^[24-26].

Knockout mice, the place a particular gene was once made inoperable by using a method referred to as gene knockout: The purpose is to be trained the operate of the gene's product or to simulate a ^[27-30] human disease:

- Fat mice, prone to weight problems as a result of a carboxypeptidase E ^[31] deficiency
- Powerful muscular mice, with a disabled myostatin gene, nicknamed "mighty mice."

Behaviour and Appereance

Laboratory mice have retained among the bodily and behavioural traits of house mice, nonetheless, as a result of many generations of man-made decision some of these characteristics now differ markedly. As a result of the massive number of traces of laboratory mice, it's impractical to comprehensively describe the appearance and behaviour of all these, nonetheless, they are described below for two of essentially the more commonly used lines ^[32-35].

A female C57BL/6 laboratory mouse:

1. C57BL/6 mice have a gloomy brown, almost black coat. They are extra touchy to noise and odours and usually tend to chunk than the more docile laboratory traces comparable to BALB/c.
2. Staff-housed C57BL/6 mice (and different traces) show barbering behaviour, wherein the dominant mouse in a cage selectively ^[36] gets rid of hair from its subordinate mates in cage. Mice that have been barbered

extensively can have giant bald patches on their bodies, usually across the head, snout, and shoulders, although barbering may just appear wherever on the physique. Each hair and vibrissae could also be eliminated. Barbering is more as a rule obvious in female mice; male mice are more likely to show dominance by means of fighting.

3. C57BL/6 has a number of uncommon traits which make it useful for some research stories but inappropriate for others: it is unusually sensitive to affliction and to bloodless and analgesic drugs is much less mighty in this stress. Not like most laboratory mouse traces, the C57BL/6 drinks alcoholic beverages voluntarily. It's extra inclined than common to morphine addiction, atherosclerosis, and age-associated hearing loss ^[37- 40].

BALB/c laboratory mice:

1. BALB/c is an, laboratory albino -bred strain from which a number of long-established sub strains are derived. With over 200 generations bred considering the fact that 1920, BALB/c mice are dispensed globally and are among the most extensively used inbred strains utilized in animal experimentation ^[41, 42].
2. BALB/c are famous for exhibiting excessive stages of nervousness and for being fairly resistant to weight loss plan-brought on atherosclerosis, making them a valuable model for cardiovascular research ^[43].
3. Male BALB/c mice are aggressive and will combat other adult males if housed collectively. Nevertheless, the BALB/Lac substrain is way more docile. Most BALB/c mice substrains have a long reproductive existence-span ^[44-46].
4. There are famous variations between special BALB/c substrains, although these are concept to be as a result of mutation instead than genetic contamination. The BALB/cWt is distinct in that three% of progeny display proper hermaphroditism ^[47, 51].

Application of Laboratory Mice in Human Disorder and Diseases

The biomedical application of laboratory mice promotes the research in ^[52-61]:

1. Cataracts
2. Diabetes
3. Obesity
4. Respiratory problems
5. Seizures
6. Deafness
7. Alzheimer's disease
8. Parkinson's disease
9. HIV and AIDs
10. Cancer
11. Cystic fibrosis
12. Muscular dystrophy
13. Spinal cord injuries
14. Hypertension
15. Heart disease

CONCLUSION

Laboratory mice are widely used in the medical research. Model mice have opened the path for the novel drug discovery in past few years. Model animals are used by the scientist for several diseases and disorder. Laboratory mice are being used in finding the treatment of Alzheimer's disease and Parkinson's disease by analysing the effect of different drug induced.

REFERENCES

1. Henkin. Short Review of Current Research on the Physiology and Pathology of Olfactory Detection. J Neurol Neurophysiol. 2016;7:363.

2. Yang H, et al. Generating genetically modified mice using CRISPR/Cas-mediated genome engineering. *Nat Protoc.* 2014;9:1956-68.
3. Brownstein DG et al. Pathogenesis of infection with a virulent allotropic variant of minute virus of mice and regulation by host genotype. *Lab Invest.* 1991;65:357–364.
4. Cheetham SA et al. Limited variation in the major urinary proteins of laboratory mice. *Physiol Behav.* 2009;96:253–261.
5. Suzuki N, et al. Cellular Transplantation as the Treatment of Alzheimer's disease in Mouse Models. *J Alzheimers Dis Parkinsonism.* 2016;6:219.
6. Maria L and Björn O. Lessons Learned from Transgenic Mouse Models for the Therapeutic Use of Drp1 Inhibitors. *Single Cell Biol.* 2016;5:135.
7. Mayordomo-Aranda E et al. Human Angiosarcoma: A Histological and Biological Phenotyping Using Xenografts in Nude Mice: Analysis of Five Cases. *J Clin Exp Pathol.* 2015;5:213.
8. Berry RJ and Bronson FH. Life history and bioeconomy of the house mouse. *Biol Rev.* 1992; 67:519–550.
9. Pillai and Shiv. "History of Immunology at Harvard". Harvard Medical School:About us. Harvard Medical School. Retrieved 19 December 2013.
10. Liu X et al. Evaluation of AAVMediated Gene Therapy with Reduced Vector Volume in Cngb3 Knockout Mice, a Model of Achromatopsia. *Hereditary Genet.* 2016;5:163.
11. Liu BL et al. ICP34.5 deleted herpes simplex virus with enhanced oncolytic, immune stimulating, and anti-tumour properties. *Gene Ther.* 2003;10:292-303.
12. Aguirre-Rueda D et al. Pro-Oxidant and Inflammatory Mediators Produced In Transgenic Mice (App/Ps1). *Biochem Physiol.* 2013;2:121
13. Biswas K and Sharan SK. Manipulating the Mouse Genome Using Recombineering. *Adv Genet Eng.* 2013;2:108.
14. Church DM et al. modernizing reference genome assemblies. *PLoS Biol.* 2011;9:e1001091.
15. Keane TM et al. Structural variation in mouse genomes. *Front Genet.* 2014;5:192
16. Waterston RH et al. Initial sequencing and comparative analysis of the mouse genome. *Nature.* 2002;420:520–562.
17. Churchill GA et al. The Diversity Outbred mouse population. *Mamm Genome.* 2012; 23:713–718.
18. Eppig JT et al. The Mouse Genome Database (MGD): facilitating mouse as a model for human biology and disease. *Nucleic Acids Res.* 2015;43: 726–736.
19. Fahey et al. *Mamm Genome.* 2013;24: 89.
20. Li H et al. Application of APP/PS1 Transgenic Mouse Model for Alzheimer's Disease. *J Alzheimers Dis Parkinsonism.* 2015;5:201.
21. Kumar V et al. CD4+T Cells Expansion in P. berghei (NK-65) Infected and Immunized BALB/C Mice. *J Clin Exp Pathol.* 2015;5:229.
22. Moreno-Ortiz H et al. Accumulation of Microsatellite Instability in Superoxide Dismutase-1 Knockout Mice: A Possible Predictor of Germ Cell Development or Tumorigenesis. *JFIV Reprod Med Genet.* 2015;3: 144.
23. Stricker et al. The regulation of body temperature in rats and mice in heat: Effects of desalivation and the presence of a water bath. *Commun. Behav. Biol.* 1968;2: 113–119.
24. Zeiler FA. NMDA Receptor Antagonism in Refractory Status Epilepticus: Right Idea, Wrong Target? *Brain Disord Ther.* 2015;4:195.
25. Werner FM and Covenas R. Symptoms and Therapeutic Options of the Anti-NMDA Receptor Encephalitis According To a Neural Network. *Anat Physiol.* 2016;6: 136.
26. Zhou LL. GluN2B-NMDA Receptors in Alzheimer's disease: What Do they got to do with AD? *J Neurol Disord.* 2015;3: 118.
27. Patel BD et al. Quantification of Newer Anti-Cancer Drug Clofarabine in their Bulk and Pharmaceutical Dosage Form. *J Chromatogr Sep Tech.* 2016;7:328.
28. Xiao X et al. 1H NMR Metabolomics Study of Spleen from C57BL/6 Mice Exposed to Gamma Radiation. *Metabolomics.* 2016;6:165.
29. Steensma et al. "Abbie Lathrop, the "Mouse Woman of Granby": Rodent Fancier and Accidental Genetics Pioneer". *Mayo Clinic Proceedings.* Mayo Foundation for Medical Education and Research. 2010;85.
30. McReynolds et al. Open-field behavior in mice: Effect of test illumination. *Psychonomic Science.* 1967;9: 277–8.

31. Stehle F et al. Snap-shot of Serine Carboxypeptidase-like Acyltransferase Evolution: The Loss of Conserved Disulphide Bridge is Responsible for the Completion of Neo-functionalization. *J Phylogen Evolution Biol.* 2013;1:115.
32. Hegmann et al. Open-field behavior in mice: Genetic analysis of repeated measures. *Psychonomic Science.* 1968;13, 27–8.
33. DeFries et al. Genetic analysis of openfield behavior. In Lindzey, G. and Thiessen, D. D. (Eds.) *Contributions to behavior-genetic analysis: The mouse as a prototype.*
34. Berry RJ. Population dynamics of the house mouse. *Symp Zool Soc Lond.* 1981;47:395–425.
35. Collins J et al. Intestinal enzyme profiles in normal and rotavirus-infected mice. *J Pediatr Gastroenterol Nutr.* 1988;7:264–272.
36. Wang X et al. Metastatic Melanoma Induced Metabolic Changes in C57BL/6J Mouse Stomach Measured by ¹H NMR Spectroscopy. *Metabolomics.* 2014;4:135.
37. Kim SY et al. The Hypolipidemic and Hypoglycemic Activities of Fermented Brown Rice Fibers by Regulating PPARα and ChREBP in the Livers of C57BL/6J Mice. 2014.
38. Sung YH et al. Mouse genetics: catalogue and scissors. *BMB Rep.* 2012;45:686–692.
39. Accurso FJ et al. Effect of vx-770 in persons with cystic fibrosis and the g551d-cftr mutation. *The New England journal of medicine.* 2010;363: 1991-2003.
40. Pudroma X and Gongsangduoji. Comparison of 5-Aminolaevulinic Acid and its Heptyl Ester-induced Protoporphyrin IX and its Photobleaching in Human Adenocarcinoma WiDr Cells and in Athymic Nude Mice Healthy Skin. *J Nucl Med Radiat Ther.* 2012;3:131.
41. Zandieh M et al. Assessment of Protection Induced by DNA and Live Vaccine Encoding Leishmania MHC Class I Restricted Epitopes against L. major Challenge in Balb/c Mice Model. *J Microb Biochem Technol.* 2015;7:427-438.
42. Mohamed A. An Additional risk of Lung Cancer from Recurrent Exposure to Ethyl Carbamate (EC) in BALB/C Mice. *J Cancer Sci Ther.* 2015;7:359-362.
43. Saunders CJ et al. Loss of function variants in human PNPLA8 encoding calcium-independent phospholipase A2 γ recapitulates the mitochondriopathy of the homologous null mouse. *Hum Mutat.* 2015;36:301–306.
44. Huang YH et al. Nanoparticle-delivered suicide gene therapy effectively reduces ovarian tumor burden in mice. *Cancer Res.* 2009;69:6184-91.
45. Ko HJ. Recent Update of Nanobiosensors Using Olfactory Sensing Elements and Nanomaterials. *Biosens J.* 2015;4:129.
46. Brisken C and O'Malley B. Hormone action in the mammary gland. In: Bissell MJ, Polyak K, Rosen JM .*The mammary gland as an experimental model.* Cold Spring Harb. *Perspect. Biol.* Cold Spring Harbor Laboratory Press, Cold Spring Harbor. 2011; 71:85.
47. Kumar V et al. CD4+T Cells Expansion in P. berghei (NK-65) Infected and Immunized BALB/C Mice. *J Clin Exp Pathol.* 2015 5:22.
48. Latifynia A et al. Th1, Th2 Serum Cytokines and Spleen White Pulp Changes Against Preliminary L. Major Vaccine Injection and Challenge With Live L. Major Promastigotes in Balb/C Mice. *J Clin Cell Immunol.* 2015;5:281.
49. Baselga J et al. Recombinant humanized anti-HER2 antibody (Herceptin) enhances the antitumor activity of paclitaxel and doxorubicin against HER2/neu overexpressing human breast cancer xenografts. *Cancer Res.* 1998;58:2825-31.
50. Ould-Ali D et al. Bleomycin-induced Scleroderma in Nude Mice can be reversed by Injection of Adipose Tissue: Evidence for a Novel Therapeutic Intervention in Systemic Sclerosis. *J Clin Exp Dermatol Res.* 2012;3:164.
51. Mayordomo-Aranda E, et al. Human
52. Sugama S, et al .Effect of Chronic Stress in the Onset of Parkinson's Disease: Possible Role of Microglial Cells in Neuroinflammation. *J Neurol Disord.* 2015;S2:001
53. Biswas A and Das SK. Alzheimer and Parkinson's disease -Two Faces of the Same Disease? *J Alzheimers Dis Parkinsonism.* 2016;6:222.
54. Melamed I. Alzheimer's disease of the Immune System: A New Variant of Immune Deficiency. *Immunother Open Acc.* 2016;2:115.
55. Xenos M and Raptis A. Fluid Structure Interaction for Biomedical Applications. *J Appl Computat Math.* 2014;3:187

56. Angiosarcoma A. Histological and Biological Phenotyping Using Xenografts in Nude Mice: Analysis of Five Cases. *J Clin Exp Hamzawy MA, Abo-youssef AM, Salem HF, Sameh 23.*
57. Christensen HR, et al. Estrogen regulation of the dopamine-activated GIRK channel in pituitary lactotrophs: implications for regulation of prolactin release during the estrous cycle. *Am J Physiol Regul Integr Comp Physiol.* 2011;301:R746.
58. Jaini R et al. An autoimmune-mediated strategy for prophylactic breast cancer vaccination. *Nat Med.* 2010;16:799-803.
59. McPhail GL et al: The first therapy acting on the primary cause of cystic fibrosis. *Drugs of today.* 2013;49: 253-260.
60. Sauvageau M et al. Multiple knockout mouse models reveal lincRNAs are required for life and brain development. *ELife.* 2013;2:e01749.
61. Gosling et al. *Behav Ecol Sociobiol* 2000;48: 328.