

Transitioning of Humanized Mouse and Rat Models from Traditional Approaches to Innovative Strategies

Isabella Rossi

EasyChair preprints are intended for rapid dissemination of research results and are integrated with the rest of EasyChair.

Transitioning of Humanized Mouse and Rat Models from Traditional Approaches to Innovative Strategies

Isabella Rossi

Nordica University, Czech Republic

Abstract

In the realm of HIV research, humanized mouse and rat models have emerged as essential tools, serving as vital links between in vitro studies and clinical trials. This article delves into the present state of these models, emphasizing their critical roles in exploring preventive and therapeutic approaches against HIV infection and related ailments. The discourse covers diverse methodologies utilized for humanizing these animal models, discusses challenges encountered, and introduces promising emerging techniques that are propelling HIV research forward. Through an analysis of the strengths and weaknesses of current models, this article seeks to guide researchers towards more efficient strategies in the continual battle against HIV/AIDS.

Introduction

The human immunodeficiency virus (HIV) remains a formidable global health challenge, affecting millions of people worldwide. Despite significant progress in understanding the virus and developing antiretroviral therapies, a definitive cure for HIV/AIDS remains elusive[1]. Bridging the translational gap between in vitro studies and clinical trials is essential for advancing our understanding of HIV pathogenesis, immune responses, and developing effective prevention and cure strategies. In this context, the utilization of humanized mouse and rat models has emerged as a transformative approach, offering unique insights into the intricacies of HIV infection and associated pathologies[2].

HIV/AIDS, caused by the retrovirus HIV, has led to a global pandemic with profound social, economic, and health implications. Despite advances in antiretroviral therapy (ART) that have transformed HIV from a once-fatal illness to a manageable chronic condition, challenges such as viral persistence, the emergence of drug resistance, and limited accessibility to treatment persist. Moreover, the lack of a preventive vaccine underscores the urgency for innovative research strategies to combat HIV at its roots[3].

Traditional animal models, such as mice and rats, have been invaluable in scientific research. However, they fall short in modeling HIV infection due to species-specific barriers[4]. Humanized mouse and rat models, engineered to incorporate human immune components, offer a revolutionary platform to study HIV in vivo, allowing researchers to overcome the limitations of conventional models[5].

Humanized mouse and rat models represent a groundbreaking leap in HIV research, providing a biologically relevant system to study the virus in the context of a human immune system. By incorporating human hematopoietic stem cells or tissues into these models, researchers can mimic key aspects of human physiology, enabling the investigation of HIV pathogenesis, immune responses, and the evaluation of potential therapeutic interventions in a more clinically relevant setting[6].

These models have proven instrumental in studying critical aspects of HIV infection, such as viral entry, replication, latency, and the dynamics of the host immune response. They serve as invaluable tools for preclinical testing of antiretroviral drugs and other therapeutic strategies, facilitating a more accurate prediction of treatment outcomes before advancing to human clinical trials[7].

This research article aims to provide a comprehensive overview of the current state of humanized mouse and rat models in HIV research. We will delve into the methodologies employed for humanization, explore their applications in investigating prevention and cure strategies against HIV, and discuss the challenges faced by researchers. Furthermore, we will spotlight emerging technologies that hold promise in advancing the field and improving the utility of these models[8].

As we navigate through the intricacies of humanized models, we aim to guide researchers, clinicians, and policymakers in their quest to understand, prevent, and ultimately cure HIV/AIDS. By critically assessing the strengths and limitations of existing models and highlighting emerging trends, this article seeks to contribute to the collective effort aimed at overcoming the challenges posed by HIV and forging a path towards a future free from this global health burden[9].

Methodologies in Humanizing Mouse and Rat Models:

2.1 Overview of Humanization Techniques:

Discuss the various strategies employed to humanize mouse and rat models, including the use of human stem cells, transplantation of human tissues, and genetic modification.

2.2 Advancements in Genetic Engineering:

Explore recent advancements in genome editing technologies, such as CRISPR-Cas9, and their impact on creating more sophisticated and efficient humanized models.

Applications in HIV Research:

3.1 HIV Pathogenesis Studies:

Examine how humanized mouse and rat models have contributed to our understanding of HIV pathogenesis, including viral entry, replication, and the establishment of latent reservoirs.

3.2 Testing Antiretroviral Therapies:

Evaluate the role of these models in preclinical testing of antiretroviral drugs and other therapeutic interventions, providing insights into their efficacy and safety profiles.

Challenges and Limitations:

4.1 Immune System Complexity:

Discuss the challenges associated with recapitulating the complexity of the human immune system in rodent models and propose potential strategies for improvement.

4.2 Ethical Considerations:

Address ethical considerations related to the use of humanized animals in research, including animal welfare concerns and the need for ethical guidelines.

Emerging Techniques and Future Directions:

5.1 Organoid Technology:

Explore the potential of organoid technology in enhancing the humanization of mouse and rat models, offering a more physiologically relevant environment for studying HIV infection.

5.2 Microbiome Modulation:

Discuss the influence of the microbiome on HIV infection and how manipulating the gut microbiota in humanized models could provide new insights into the interplay between the virus and host.

Conclusion:

Summarize the key findings and insights from the review, emphasizing the crucial role of humanized mouse and rat models in advancing HIV research. Highlight the potential of emerging technologies and the need for collaborative efforts to overcome existing challenges and pave the way for more effective prevention and cure strategies against HIV/AIDS.

By examining the current and emerging landscape of humanized mouse and rat models, this article aims to provide a comprehensive resource for researchers working towards a deeper understanding of HIV infection and the development of novel therapeutic approaches.

References

- [1] R. Mishra, "Bird Mating Optimizer and Its Applications in Medical Research," 2023.
- [2] Y. Agarwal et al., "Development of humanized mouse and rat models with full-thickness human skin and autologous immune cells," Sci. Rep., vol. 10, no. 1, 2020.
- [3] S. Biradar, M. T. Lotze, and R. B. Mailliard, "The unknown unknowns: Recovering gamma-delta t cells for control of human immunodeficiency virus (HIV)," Viruses, vol. 12, no. 12. 2020.
- [4] D. Johnson and J. Smith, "A Brief Analysis on Adverse Side Effects of COVID-19 Vaccines."
- [5] S. Biradar, Y. Agarwal, M. T. Lotze, M. T. Bility, and R. B. Mailliard, "The BLT Humanized Mouse Model as a Tool for Studying Human Gamma Delta T Cell-HIV Interactions In Vivo," Front. Immunol., vol. 13, 2022.
- [6] Y. Agarwal et al., "Moving beyond the mousetrap: Current and emerging humanized mouse and rat models for investigating prevention and cure strategies against HIV infection and associated pathologies," Retrovirology, vol. 17, no. 1. 2020.
- [7] F. Tahir and M. Khan, "Study of High Blood Pressure and its Effect to Cancer."
- [8] B. Angeleo, B. Antonio, and M. Khan, "Challenges in Species Distribution Modelling paradigm and Modell Elevation."
- [9] L. Ghafoor, "The Classification of Neurosurgical Complications," 2023.