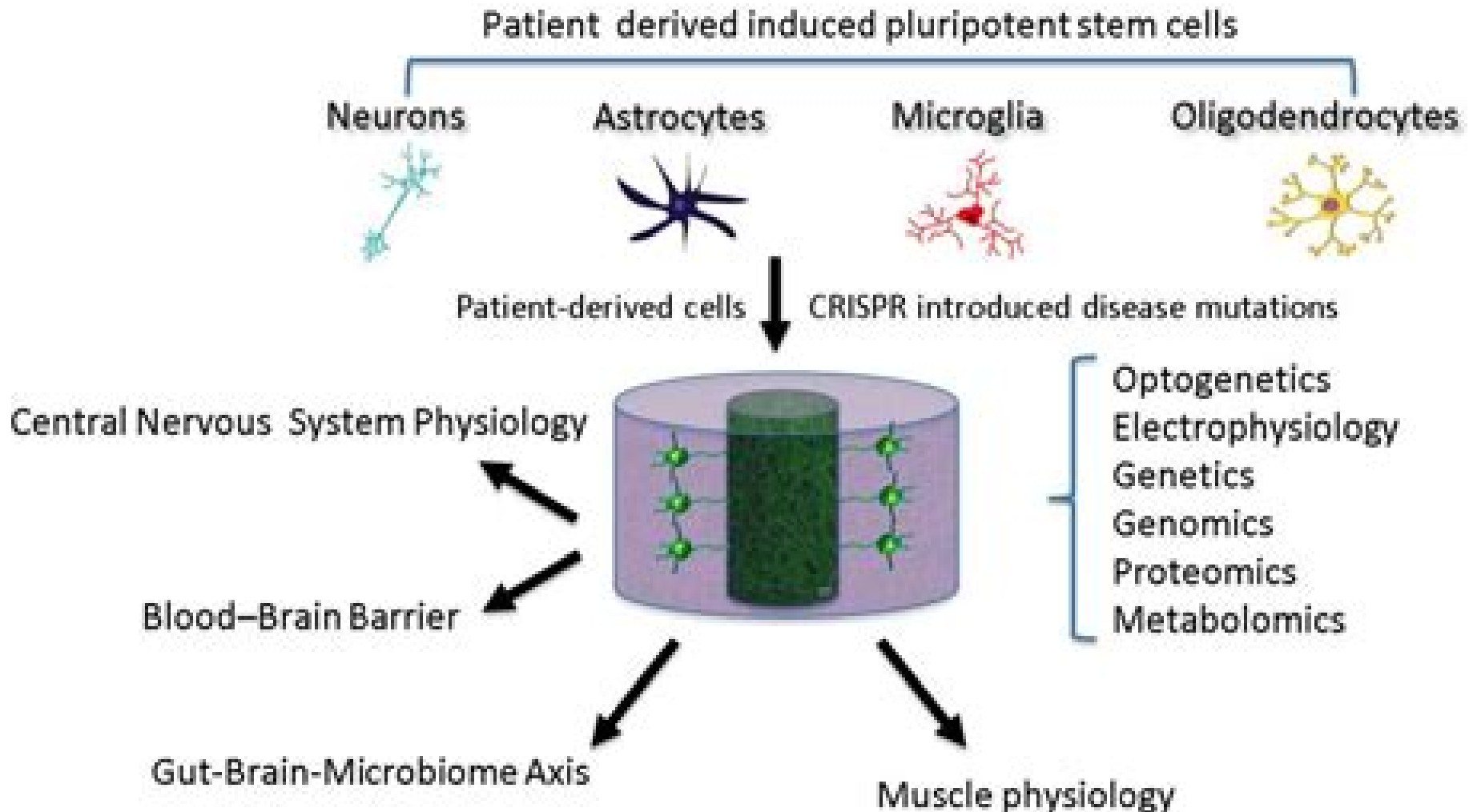


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Understanding how disease mutations may cause neurological defects



Models Of Human Neurological Diseases

Ebru Emekli-Alturfan



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Models of human neurological diseases, 1971 *Models of Human Neurological Diseases* Harold L. Klawans, 1974

New Animal Models of Human Neurological Diseases P. Poindron, P. Piguet, 2008-02-25 This publication describes in detail some animal models of outstanding importance for the study of human neurological diseases An introductory article provides a definition of animal models looks at the history and ethics of their use and highlights the reasons why animal models are so useful in scientific research The following chapters present some of the newer models which play a key role in the study of amyotrophic lateral sclerosis multiple sclerosis stroke and peripheral neuropathy Their specific features are detailed in the method section of each paper and their application to the study of human neurological diseases is discussed in a comprehensive manner i e limits and challenges posed by these models are evaluated and potential improvements suggested A valuable single source of information this book is recommended reading for neuroscientists as well as for neurologists and neurosurgeons interested in preclinical research on human neurological diseases Neural Dynamics of Neurological Disease Christopher A. Shaw, 2017-04-17 The emerging understanding of age related neurological disorders suggests that notions of a single causal gene toxin being responsible is likely incorrect Neurological disorders probably arise due to a unique intersection of multiple genetic and toxic factors combined with additional contributions of age stage of development immune system actions and more This perspective leads to the view that rather than reflecting only one pathway to end state disease each is a spectrum disorder and every individual case is therefore unique Neural Dynamics of Neurological Disease argues for a fundamental rethinking of what we think we know about neurological disorders how they arise and progress and crucially what might be done to cure them It first introduces the concept of neural dynamics of neurological disease then examines various diseases and gives examples of the interplay of elements such as neural systems cell types and biochemical pathways that can contribute to disease The concluding chapters point the way to how the emerging notion of neurological disease as a dynamic process may lead to more successful treatment options Providing a cross disciplinary approach to understanding the origin and progression of neurological disease Neural Dynamics of Neurological Disease is a timely and valuable resource for neuroscientists researchers and clinicians **Animal Models of Human Disease** Sara Green, 2024-05-08 The crucial role of animal models in biomedical research calls for philosophical investigation of how and whether knowledge about human diseases can be gained by studying other species This Element delves into the selection and construction of animal models to serve as preclinical substitutes for human patients It explores the multifaceted roles animal models fulfil in translational research and how the boundaries between humans and animals are negotiated in this process The book also covers persistent translational challenges that have sparked debates across scientific philosophical and public arenas regarding the limitations and future of animal models Among the are persistent tensions between standardization and variation in medicine as well as between strategies aiming to reduce and recapitulate

biological complexity Finally the book examines the prospects of replacing animal models with animal free methods The Element demonstrates why animal modeling should be of interest to philosophers social scientists and scientists alike

Peroxisome Biology: Experimental Models, Peroxisomal Disorders and Neurological Diseases Gérard

Lizard,2021-01-08 This book provides an overview of the biology and biochemistry of peroxisomes and discusses the contribution of these organelles to peroxisomal and neurodegenerative diseases It begins with a detailed introduction to the biogenesis and metabolic functions of peroxisomes and highlights their role in oxidative stress and in lipid metabolism such as fatty acid oxidation The following chapters focus on the molecular and clinical aspects of peroxisomal disorders caused by defects in peroxisomal function In particular the biological aspects of peroxisomal biogenesis disorders such as Zellweger syndrome and Heimler syndrome are discussed This includes their underlying genetic causes as well as the biochemical and metabolic defects associated with the disorders In addition several chapters cover recent observations suggesting an association between peroxisomal dysfunction and neurodegenerative diseases such as Alzheimer s Multiple Sclerosis and other degenerative cerebellar pathologies The final section of the book discusses important cell and animal models for studying the role of peroxisomes in human diseases and presents current therapeutic strategies for their treatment This book deals with a highly topical subject that is at the heart of current research and represents a valuable contribution for all students and researchers who want to understand the complex biology of peroxisomes and their role in human diseases

Toxin-Induced Models of Neurological Disorders A.J. Nonneman,M.L. Woodruff,2013-11-11 This volume discusses and evaluates the use of neurotoxins to produce animal models of human neurodegenerative diseases The models presented use neuro toxins to induce some or most of the pathophysiological changes including behavioral that characterize the most studied motor neuron disorders and cognitive disorders dementia in humans Within the biomedical research community there is little doubt about the utility of such models But with increasing frequency individuals concerned with animal rights question the validity of such models and argue that they actually produce no useful information for understanding or developing treatments for human disorders Each of the chapters will address this issue considering the utility validity generalizability and limitations of the models presented Chapter 1 begins with a brief review of the ethics of animal use in neuro biological research including a discussion of the criteria that may be used to evaluate animal models of human disease and extrapolate from the model to appropriate questions regarding humans The limitations of such extrapolation are also discussed with special consideration of issues specific to the use of neurotoxins Chapter 2 extends this theme with further consideration of issues and strategies involved in developing neurotoxin induced models of neuro degenerative disorders and assessing risk factors associated with neurotoxin exposure It then narrows to evaluate the use of a model systems approach to explore neurotoxin induced learning and memory deficits in animals as related to humans with dementia *Mouse Models in the Study of Genetic Neurological Disorders* Brian Popko,2012-12-06 The number of mouse models that are

available for the study of human genetic neurological disorders is large and growing rapidly Therefore it was difficult to select the models that were reviewed in this volume Clearly there are important models that are not discussed and perhaps a volume twice this size would have been more appropriate Moreover the pace at which new models are being developed and analyzed is rapid As this volume goes to press I am sure that additional mouse genes responsible for naturally occurring neurological disorders are being discovered and that many new transgenic and mutant mouse strains are being developed Therefore this volume should not be viewed as a comprehensive compendium but rather as an update of work in progress It is exhilarating to witness the fast pace at which these models are being established as important tools in the study of basic neuroscience and neurological disorders It will be even more exciting to see their utilization in the development and testing of therapeutic interventions for these diseases I would like to thank each of the authors who have contributed to this volume for their time and their expertise I would also like to thank Drs Timothy Coetzee and Joshua Corbin for their advice in the selection of the topics covered I am deeply indebted to Dr Kunihiko Suzuki who first approached me with the idea for this volume for his guidance throughout its preparation *Molecular and Functional Models in Neuropsychiatry* Jim J.

Hagan,2011-04-02 The development of more effective treatments for neuropsychiatric disorders requires scientific progress on a broad front Animal models have a vital role to play in advancing the field When deployed in conjunction with detailed study of these diseases in man they bring the power to make controlled experimental interventions which allow the functional consequences of genetic variations and polymorphisms to be understood in terms of their cellular systems and behavioural effects Further they provide a means by which complex cognitive and behavioural phenomena may be dissected and understood Finally they provide a bridge to understanding the effects of drugs on the functioning of the central nervous system thereby improving our understanding of the actions of those drugs in man **Zebrafish Models of**

Neurodegenerative Disorders Ebru Emekli-Alturfan,2025-09-08 Advancing our understanding of neurodegenerative diseases demands innovative tools and approaches Zebrafish Models in Neurodegenerative Disorders offers a comprehensive examination of zebrafish as a translational model for studying the molecular and cellular mechanisms of diseases such as Alzheimer s Parkinson s and Huntington s diseases Beginning with the first section informing about the basic molecular mechanisms of neurodegenerative disorders biomarkers and therapeutic approaches the book continues with the detailed exploration of zebrafish physiology genetics and behavioural phenotypes as they pertain to neurodegenerative pathologies insight into cutting edge technologies including CRISPR Cas9 genome editing and in depth analysis of zebrafish models highlighting their strengths in modelling disease progression and therapeutic testing

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