G. Lubec (ed.)

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manipulation techniques new interest has developed in the biological mechanisms of aggression The primary goal of this title is to summarise and synthesis recent advances in the subject **Emerging Mechanisms in Neuronal Signaling: From** Cell Biology to Pathogenesis Mario Eduardo Guido, Gabriela Alejandra Salvador, Alejandra Alonso, 2020-11-12 This eBook is a collection of articles from a Frontiers Research Topic Frontiers Research Topics are very popular trademarks of the Frontiers Journals Series they are collections of at least ten articles all centered on a particular subject With their unique mix of varied contributions from Original Research to Review Articles Frontiers Research Topics unify the most influential researchers the latest key findings and historical advances in a hot research area Find out more on how to host your own Frontiers Research Topic or contribute to one as an author by contacting the Frontiers Editorial Office frontiers in org about International Review of Research in Mental Retardation, 2011-09-06 International Review of Research in Mental Retardation is an ongoing scholarly look at research into the causes effects classification systems syndromes etc of mental retardation Contributors come from wide ranging perspectives including genetics psychology education and other health and behavioral sciences Volume 36 of the series offers chapters on newborn screening for intellectual disability responsive parenting Trisomy 21 Alzheimer's disease in adults with Down Syndrome foolish action in adults with intellectual disabilities animal models of self injurious behavior theoretical and methodological issues in sibling research understanding individual differences in adaptation in parents of children with intellectual disabilitie and What Do You Think If using vignettes to study attitudes toward adult sibling caregiving and competence of parents of children with disabilities. The wide range of topics covered in these chapters make Volume 36 of the International Review of Research in Mental Retardation a particularly valuable resource for academic researchers in developmental and cognitive psychology as well as those in neuropsychology Provides the most recent scholarly research in the study of mental retardation A vast range of perspectives is offered and many topics are covered An excellent resource for academic researchers The Molecular Biology of Down Syndrome G. Lubec, 1999-12-02 This book contains updated reviews and original research work on Down Syndrome focussing on brandnew results in neurobiology in particular results on gene hunting subtractive hybridization differential display and neurochemistry The book provides new data such as a subtractive library of Down Syndrome brain showing cDNAs that are overexpressed or downregulated and can be regarded as a source for further research on the preliminary transcriptional data given A 2D electrophoretic map of human brain proteins including Down Syndrome brain protein expression established by in gel digestion of spots with subsequent MALDI identification provides the scientific basis for protein work to the neuroscientist Altogether the book provides a series of new candidate genes possibly involved in Down Syndrome neurobiology tools for neuroscience studies on Down Syndrome brain thus serving as a manual and updated views and aspects on Down Syndrome pathobiology Oesterreichische Bibliographie, 2000 Advances in Down Syndrome Research Gert Lubec, 2003-12-19 Advances in Down Syndrome Research represents updated research in several areas of Down Syndrome DS A new promising

animal model of DS is reported and this opens new opportunities to study pathomechanisms and pharmacological approaches as it is more than difficult to carry out studies in humans and the clinical features are highly variable In terms of biology cell cycle and stem cell studies and in terms of biochemistry relevance of studies on a specific protein kinase channels transporters superoxide dismutase antioxidant system chromosome assembly factor and other important biological structures are provided And again the gene dosage hypothesis is addressed and although the vast majority of chromosome 21 gene products is unchanged in fetal DS brain a few specific chromosome 21 encoded structures including transcription factors are indeed overexpressed although findings in fetal DS are different from those in adult DS brain when Alzheimer like neuropathology supervenes Protein Expression in Down Syndrome Brain G. Lubec, 2013-12-01 When we worked on Down Syndrome brain in the past we have been focus ing on adult brain This was a major step forwards as most work on Down Syndrome was carried out on fibroblasts or other tissues and moreover we introduced proteomics to identify and quantify brain protein expression We considered evaluation of brain protein expression in Down Syndrome brain by and by more important than gene hunting at the nucleic acid level realiz ing the long unpredictable way from RNA to protein The availability of fetal samples along with the proteomic appproach stimulated and reinforced studies on Down Syndrome brain And indeed it was found out that some observations on aberrant protein expression in adult Down Syndrome brain could not be verified in the fetal samples indicating that neurodegeneration in adult Down Syndrome brain may have been responsible rather than trisomy 21 Using brains from the early second trimester of gestation led to the generation of a series of clues for the under standing of aberrant wiring of the brain in Down Syndrome and enabled the determination of altered key functions in early life e g undetectably low drebrin was observed in Down Syndrome cortex an integral constituent and marker for dendritic spines main effectors of cross talk between neurons In addition evaluation of the nature of the neuronal deficits in terms of neuro transmission markers could be established as well as neuronal density in fetal Down Syndrome cortex

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discussion of the genetic models of down syndrome across diverse species including rodents zebrafish drosophila and human organoids Authors use a cross cutting approach to compare the strengths and weaknesses of each model system explore how to use model organisms to study human diseases and capture the status of the field This book is a useful resource for biomedical researchers and students interested in using model systems to study Down syndrome and learn about Down syndrome and other chromosomal disorders

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