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because may occur at young ages The only known cause of the disease is associated with genetic mutations mainly in the gene encoding superoxide dismutase 1 familial ALS whereas there is no known cause of the sporadic form of ALS SALS which comprises 90% of cases Both ALS types develop similar histopathological and clinical characteristics and there is no treatment or prevention of the disease Because effective treatments for ALS as for other neurodegenerative diseases can only result from the knowledge of their cellular and molecular pathophysiological mechanisms research on such mechanisms is essential Although progress in neurochemical physiological and clinical investigations in the last decades has identified several mechanisms that seem to be involved in the cell death process such as glutamate mediated excitotoxicity alterations of inhibitory circuits inflammatory events axonal transport deficits oxidative stress mitochondrial dysfunction and energy failure the understanding of the origin and temporal progress of the disease is still incomplete and insufficient Clearly there is a need of further experimental models and approaches to discern the importance of such mechanisms and to discover the factors that determine the selective death of motor neurons characteristic of ALS in contrast to other neurodegenerative diseases such as Parkinson's and Alzheimer's disease Whereas studies in vitro in cell cultures tissue slices or organotypic preparations can give useful information regarding cellular and molecular mechanisms the experiments in living animal models obviously reflect more closely the situation in the human disease provided that the symptoms and their development during time mimics as close as possible those of the human disease It is necessary to correlate the experimental findings in vitro with those in vivo as well as those obtained in genetic models with those in non genetic models aiming at designing and testing therapeutic strategies based on the results obtained Issues in Materials and Manufacturing Research: 2012 Edition, 2013-01-10 Issues in Materials and Manufacturing Research 2012 Edition is a ScholarlyEditions eBook that delivers timely authoritative and comprehensive information about Molecular Modeling The editors have built Issues in Materials and Manufacturing Research 2012 Edition on the vast information databases of ScholarlyNews You can expect the information about Molecular Modeling in this eBook to be deeper than what you can access anywhere else as well as consistently reliable authoritative informed and relevant The content of Issues in Materials and Manufacturing Research 2012 Edition has been produced by the world's leading scientists engineers analysts research institutions and companies All of the content is from peer reviewed sources and all of it is written assembled and edited by the editors at ScholarlyEditions and available exclusively from us You now have a source you can cite with authority confidence and credibility More information is available at http www ScholarlyEditions com American Journal of Diseases of Children ,1923 California and Western Medicine ,1926 The Official Bulletin of the Medical Women's Club of Chicago Medical Women's Club of Chicago, 1927 List of members in most numbers Neuromuscular Disorders: Management and Treatment E-Book Tulio E. Bertorini, 2010-09-08 Neuromuscular Disorders presents a multi disciplinary approach to the management and therapeutic treatment of the full range of neuromuscular disorders and resulting complications Dr Tulio Bertorini and a contributing

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