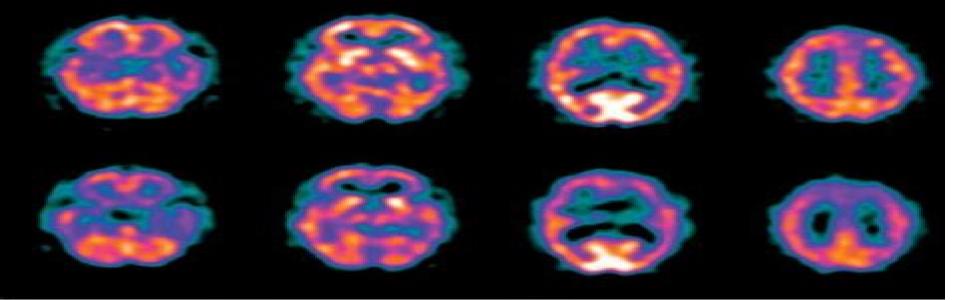
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AMYOTROPHIC LATERAL SCLEROSIS AND THE FRONTOTEMPORAL DEMENTIAS

MICHAEL J. STRONG

Serge Gauthier, Pedro Rosa-Neto

Amyotrophic Lateral Sclerosis and the Frontotemporal Dementias Michael J. Strong,2012-10-11 Over the past ten years there has been an increasing recognition that syndromes of frontotemporal dysfunction FTD are a common occurrence in patients with amyotrophic lateral sclerosis ALS Such syndromes may be present in as many as 60% of patients with ALS Conversely the occurrence of motor neuron dysfunction in patients with clinically pure frontotemporal dementia is increasingly recognized This suggests that to some extent there are overlapping syndromes in which both ALS and FTD occur within the same individual This volume summarizes the advances in our understanding of these two disorders as well as the potential relationship between the two Key topics include advances in our ability to clinically describe the frontotemporal syndromes preclinical detection neuroimaging and genetics The exploding field of new markers in neuropathology is examined as is the role of new genetic mutations in DNA RNA transport systems This book is the essential reference text for this topic and will be of interest to neurologists and neurological trainees with a clinical or research interest in the FTDs or ALS neuropsychologists neuropathologists and researchers

Multifaceted Genes in Amyotrophic Lateral

Sclerosis-Frontotemporal Dementia Henry Houlden, Alan Edward Renton, Francesca Luisa Conforti, 2021-06-28

Oxytocin in Brain Health and Disease: How can it exert such pleiotropic neuromodulatory effects? Francesca Talpo, Navjot Kaur, Gerardo Rosario Biella, 2023-06-08 Oxytocin is a pituitary hormone able to produce a multitude of heterogeneous central and peripheral responses Within the central nervous system oxytocin is synthesized by the hypothalamic parvocellular neurons and released in many different brain areas where it acts as a neuromodulator It exerts pro social and anxiolytic effects by promoting attachment trust maternal bonding social affiliation and eating and metabolic functions Imbalances in the oxytocinergic system are implicated in neuropsychiatric diseases associated with altered socio emotional competence such as autism spectrum disorder clinical depression and eating disorders Also oxytocin impairments could affect memory formation persistence and they have been found in some neurodegenerative diseases such as Alzheimer s disease and in several cancers Although substantial progress has been achieved in understanding single pieces of the complex neurobiology of the oxytocinergic system the puzzle is far from being complete It is still unclear how this single neuropeptide could exert such pleiotropic neuromodulatory effects Extending the comprehension of the mechanisms underlying the complex roles of the oxytocin will be of great interest as this molecule has the potentiality to be used as a Case Studies in Dementia: Volume 1 Serge Gauthier, Pedro Rosa-Neto, 2011-04-21 Dementia is amongst drug treatment the greatest challenges facing the medical profession as the population ages Accurate diagnosis is essential as many rarer forms of the disease are treatable if recognized early This collection of case studies from around the world illustrates both common and unusual causes of dementia emphasizing clinical reasoning integrative thinking and problem solving skills Each case consists of a clinical history examination findings and special investigations followed by diagnosis and discussion The

aim is to reinforce diagnostic skills through careful analysis of individual presenting patterns and to guide treatment decisions using state of the art diagnostic classification and tools The reader will be able to distinguish patients who need reassurance closer follow up or immediate referral to specialized services Written and edited by internationally recognized experts in dementia these case studies will inform and challenge clinicians at all stages of their careers Disease in Adults Mark B. Bromberg, 2015 Motor Neuron Disease reviews new information from 1998 as it applies to all aspects of motor neuron disease Articles included use evidence based methods to ensure that the new information is solid and advances the topic The book can be used by anyone who provides any type of care to ALS patients Diagnosing Amyotrophic Lateral Sclerosis M. Turner, L. Jenkins, 2019-11-26 A diagnosis of amyotrophic lateral sclerosis also known as Lou Gehrig s disease or motor neuron disease is a progressive neurodegenerative disorder that exerts a notorious life shortening physical toll Understandably clinicians are keen to avoid a wrong diagnosis when there are such serious consequences but any delay in diagnosis can result in unnecessary and sometimes harmful interventions and prevents prompt implementation of much needed physical and emotional support Starting from the premise that ALS is not one disease but a syndrome with a spectrum of upper and lower motor neuron involvement this highly readable resource examines the causes of diagnostic delay and how to avoid them With no diagnostic test to confirm the disease no mandatory investigations and very few plausible ALS mimics the authors take a pragmatic approach to what must always be a clinical diagnosis With case presentations and teaching points to aid understanding Fast Facts ALS will give clinicians the confidence to confirm or exclude a diagnosis of ALS so that individuals facing this most challenging of conditions can receive rapid multidisciplinary support to maximize the quality of their remaining life Contents Defining the syndrome Epidemiology and pathophysiology The first symptoms Differential diagnosis Investigations Emerging diagnostic biomarkers Brain Protein Aging and Dementia Control Gen Sobue, Naruhiko Sahara, 2019-06-28 **Bradley's Neurology in Clinical Practice** E-Book Robert B. Daroff, Joseph Jankovic, John C Mazziotta, Scott L Pomeroy, 2015-10-25 Comprehensive easy to read and clinically relevant Bradley's Neurology in Clinical Practice provides the most up to date information presented by a veritable Who s Who of clinical neuroscience Its unique organization allows users to access content both by presenting symptom sign and by specific disease entities mirroring the way neurologists practice A practical straightforward style templated organization evidence based references and robust interactive content combine to make this an ideal dynamic resource for both practicing neurologists and trainees Authoritative up to date guidance from Drs Daroff Jankovic Mazziotta and Pomeroy along with more than 150 expert contributors equips you to effectively diagnose and manage the full range of neurological disorders Easy searches through an intuitive organization by both symptom and grouping of diseases mirrors the way you practice The latest advances in clinical neurogenetics brain perfusion techniques for cerebrovascular disease the relationship between neurotrauma and neurodegenerative disease management strategies for levodopa related complications in

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