



Muscle Aging, Inclusion-Body Myositis and Myopathies

EDITED BY

Valerie Askanas and W. King Engel



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Hardcover 2012 By Valerie Askanase editor

**Emma Ciafaloni, Robert C.
Griggs, Patrick F. Chinnery**



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Muscle Aging, Inclusion-Body Myositis and Myopathies Valerie Askanas, W. King Engel, 2012-02-27 Muscle weakness with ageing is almost inevitable generally beginning to manifest beyond the age of 40 and is usually unstoppable It can lead to reduced mobility increased risk of falling injury and even death But you're just getting old is not a sufficient diagnosis Specific causes of neuromuscular symptoms may explain progressive muscle weakness and should be investigated for potential treatment Muscle Ageing Inclusion Body Myositis and Myopathies explores the clinical and pathological expression of muscle weakness in aging persons Case studies demonstrate how physicians can more accurately diagnose weakening elderly patients and make better management decisions It also explores sporadic inclusion body myositis and hereditary inclusion body myopathies The former the most common progressive muscle disease in the over 50s is frequently under diagnosed and with the increasing population of aged individuals is presenting a greater challenge This disease of muscle has pathological similarities with the well known Alzheimer and Parkinson brain diseases Edited and written by a leading international cast of authors Muscle Ageing Inclusion Body Myositis and Myopathies provides a state of the art guide to ageing associated neuromuscular disorders It should be in the hands of all those involved in the care of aging and muscle weakened patients Titles of Related Interest Neuromuscular Disorders Tawil and Venance eds ISBN 978 0 470 65456 9 European Handbook of Neurological Management Vol 1 2e Gilhus Barnes Brainin eds ISBN 978 1 4051 8533 2 European Handbook of Neurological Management Vol 2 2e Gilhus Barnes Brainin eds ISBN 978 1 4051 8534 9 **Inclusion-Body Myositis and Myopathies** Valerie Askanas, Georges Serratrice, W. King Engel, 1998-04-13 Inclusion body myositis IBM is now understood to be an important degenerative muscle disease The sporadic type s IBM is probably the most common muscle disease among those ailments that strike first in adulthood particularly people over 50 The hereditary type h IBM affects younger patients This book is devoted entirely to s IBM and h IBM Contributors discuss what is understood about the basic scientific foundations of IBMs the varied aspects of the pathology of IBMs and the application of clinical treatments One particular emphasis of the book is on the hereditary aspects of IBM and genetic predispositions to the disease

Polymyositis and Dermatomyositis Marinos C. Dalakas, 2013-10-22 Polymyositis and Dermatomyositis provides extensive information regarding Polymyositis and Dermatomyositis PM DM which is described as a heterogeneous disease complex This book is divided into four sections Part I Clinical Features covers the classification of PM DM details of the clinical presentation and the disease's association with the other connective tissue disorders and malignancies Part II Etiology and Mechanisms covers advances in the immunopathology and viral etiology of PM DM along with a frequently recognized entity inclusion body myositis Part III Diagnosis and Treatment covers the histologic muscle enzyme histochemical electron microscopic and resin histology features of PM DM along with those electromyographic features that could help make a more accurate diagnosis Part IV Overview summarizes the issues that may not have been clear and

highlights differing and unsettled views or present available data This text is directed to clinicians in private practice or in academic institutions concerned with PM DM patients including neurologists rheumatologists pediatricians dermatologists physiatrists and neuromuscular investigators This book is intended as well for neuromuscular pathologists who interpret muscle biopsy specimens and electromyographers who perform EMG studies to help determine the clinical diagnosis Researchers in immunology and immunopathology of neuromuscular diseases will find discussions in this book invaluable

Evaluation and Treatment of Myopathies Emma Ciafaloni, Robert C. Griggs, Patrick F. Chinnery, 2014 Evaluation and Treatment of Myopathies provides all the essential information needed to assess the patient who has a muscle disease or who has symptoms that suggest a muscle disease It provides practical advice on how to elicit symptoms and signs and then how to proceed to establish the diagnosis and institute appropriate treatment *Myopathies, An Issue of Neurologic Clinics* Mazen Dimachkie, 2014-07-23 Congenital myopathies Muscular dystrophies Glycogen storage diseases of muscle and Idiopathic and Inflammatory myopathies are presented in this volume of Neurologic Clinics Topics include Muscle channelopathies Pompe disease Congenital myopathies and muscular dystrophies Duchenne and Becker muscular dystrophies Distal myopathies Limb girdle muscular dystrophy Fascioscapulohumeral muscular dystrophy Myotonic dystrophy Metabolic and mitochondrial myopathies Sporadic inclusion body myositis Toxic myopathies Idiopathic Inflammatory myopathies Approach to muscle disease **Myositis: New Insights for the Healthcare Professional: 2012 Edition**, 2012-12-10 Myositis New Insights for the Healthcare Professional 2012 Edition is a ScholarlyPaper that delivers timely authoritative and intensively focused information about Myositis in a compact format The editors have built Myositis New Insights for the Healthcare Professional 2012 Edition on the vast information databases of ScholarlyNews You can expect the information about Myositis 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 Myositis New Insights for the Healthcare Professional 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> **Inclusion-body myositis: clinical and pathologic aspects, and basic research potentially relevant to treatment** American Academy of Neurology, 2006 *Inclusion Body Myositis* Marinos C. Dalakas, W. King Engel, 2006

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