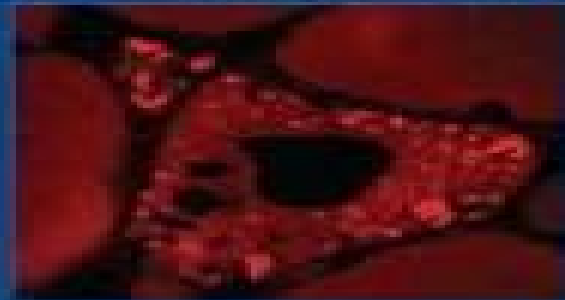


Muscle Aging, Inclusion-Body Myositis and Myopathies

EDITED BY

Valerie Askanas and W. King Engel



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Author Valerie Askanas Published On March 201

Mazen Dimachkie



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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

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

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 **Evaluation and Treatment of Myopathies** Emma

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