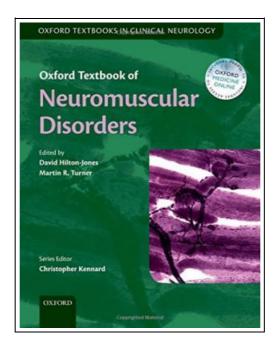
# Oxford Textbook of Neuromuscular Disorders



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## OXFORD TEXTBOOK OF NEUROMUSCULAR DISORDERS



Book Condition: New. Publisher/Verlag: Oxford University Press | Includes 12 months access to the online versionPart of the Oxford Textbooks in Clinical Neurology series, the Oxford Textbook of Neuromuscular Disorders covers the scientific basis, clinical diagnosis, and treatment of neuromuscular disorders, with a particular focus on the most clinically relevant disorders. | Part of the Oxford Textbooks in Clinical Neurology series, the Oxford Textbook of Neuromuscular Disorders covers the scientific basis, clinical diagnosis, and treatment of neuromuscular disorders with a particular focus on the most clinically relevant disorders. The book is organized into seven sections, starting with the general approach to the patient with neuromuscular disorders and then focusing on specific neuromuscular conditions affecting theperipheral nervous system from its origins at the spinal cord anterior horn on its outward course to their effector muscles and the inbound sensory pathways. Chapters on specific neuromuscular conditions are illustrated with typical case histories and their presenting features, allowing readers to put rarerconditions into their clinical context more easily. The concurrent online version allows access to the full content of the textbook, contains links from the references to primary research journal articles, allows full text searches, and provides access to figures and tables that can be downloaded to PowerPoint®. This textbook will serve as a useful reference for neurologists and specialist trainees in neurology, neuroscientists, neurophysiologists, and other healthcare professionals. | Section 1: Approach to the patient; 1 David Hilton-Jones and Martin R. Turner: Eliciting the history; 2 David Hilton-Jones and Martin R. Turner: Genetic considerations; 3 David Hilton-Jones and Martin R. Turner: Examination; Section 2: Anterior Horn; 4 Kevin Talbot and Martin R. Turner: Amyotrophic Lateral Sclerosis; 5 Dirk Baumer and Kevin Talbot: Spinal muscular atrophy and hereditary motor neuropathy; 6 Kenneth Fischbeck and Christopher Grunseich: Kennedy's disease; 7 Robin S. Howard: Poliomyelitis; Section...

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