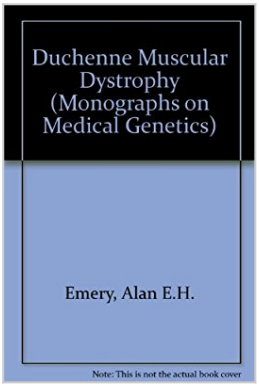

Duchenne Muscular Dystrophy (Oxford Monographs on Medical Genetics)

By Alan E. H. Emery

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Based on the author's study of some 200 patients, and on a detailed review of the literature, this volume examines all aspects of Duchenne muscular dystrophy--the second most common genetic disorder affecting humans. Topics include the history of the disease, clinical features, involvement of tissues other than muscle, laboratory investigations, differential diagnosis, biochemistry, pathogenesis, genetics, molecular pathology, prevention, genetic counselling, and surgical and medical management. The book is illustrated with numerous photographs and line drawings, and includes an extensive bibliography.

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Editorial Review

Review
'With the same fluency and flow which he achieved in the first edition, Alan Emery has succeeded in a masterly fashion in assimilating and digesting the essentials from all this new information and integrating it into this new volume. Once again he has comprehensively covered the whole field.
His vast personal experience both as a geneticist and a clinician are clearly apparent from the confidence as well as the sensitivity with which he has covered many of the topics. This book should find a wide appeal ... both as a source of reference as well as an interesting and informative read.'
Victor Dubowitz, Royal Postgraduate Medical School. London, Neuromuscular Disorders, Vol. 3, No. 2, 1993

'This book is an excellent review of the history of Duchenne muscular dystrophy ... remarkably current in the new techniques, a credit to the author. There is something for everyone interested in neuromuscular disease in this book. The book is extremely well written, and easy to read and I
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'Professor Emery's book is a useful reference text for all of those involved in the care of patients with Duchenne muscular dystrophy or research into the muscular dystrophies. This text provides a rounded view of all of the many issues involved ... a useful starting point for those who care
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