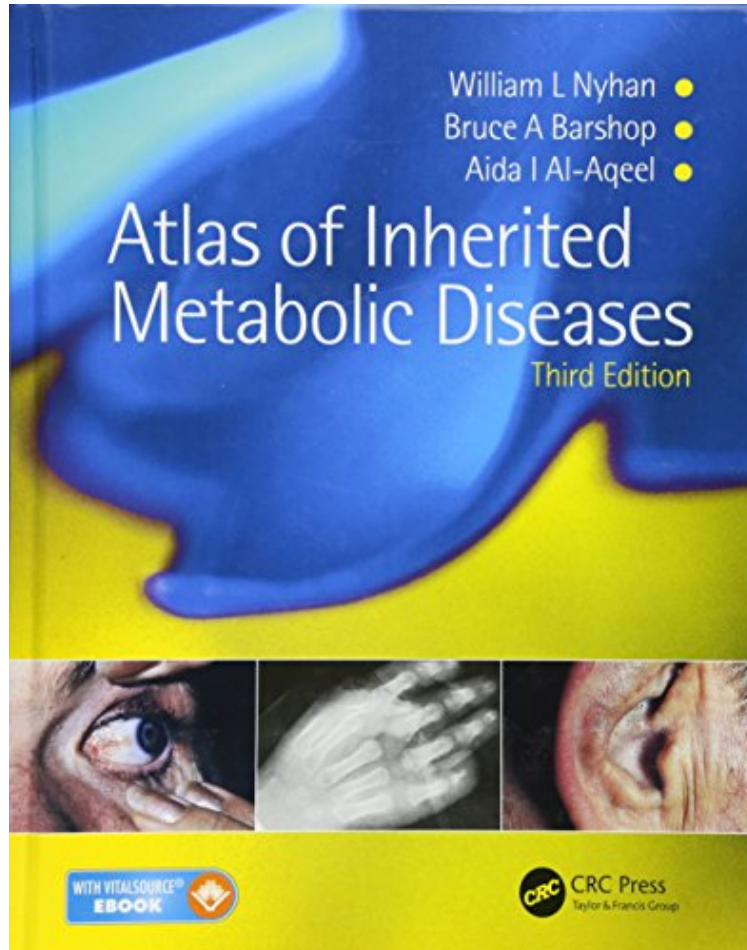


[Read free ebook] Atlas of Inherited Metabolic Diseases 3E

Atlas of Inherited Metabolic Diseases 3E

William L Nyhan, Georg F. Hoffman, Bruce A Barshop, Aida I Al-Aqeel
DOC | *audiobook | ebooks | Download PDF | ePub



[Download](#)

[Read Online](#)

#550575 in Books CRC Press 2011-12-30 Original language: English PDF # 1 8.60 x 1.70 x 10.90l, 5.82 #File Name: 1444112252888 pages | File size: 68.Mb

William L Nyhan, Georg F. Hoffman, Bruce A Barshop, Aida I Al-Aqeel : Atlas of Inherited Metabolic Diseases 3E before purchasing it in order to gage whether or not it would be worth my time, and all praised Atlas of Inherited Metabolic Diseases 3E:

0 of 0 people found the following review helpful. Great pictures!By HectorAtlas of Inherited Metabolic Diseases 3th edition is a great book! It offers you very nice clinical pictures together with the clinical, biochemical and molecular features of each disease.0 of 0 people found the following review helpful. No proper bill on !By Julia SzaboThe book is perfect. I am disappointed, as I requested a VAT bill, but you did not provide it to me!0 of 0 people found the following review helpful. Well written easy to followBy nirtakA must if you must have the information for clarifying a diagnosis. Well written easy to follow.

In a field where even experts may find that years have elapsed since they last encountered a child with a given

disorder, it is essential for the clinician to have a comprehensive source of practical and highly illustrated information covering the whole spectrum of metabolic disease to refer to. The third edition of this highly regarded book, authored by three of the foremost authorities in pediatric metabolic medicine, fulfils this need by providing an invaluable insight into the problems associated with metabolic diseases. The Atlas of Metabolic Disease is divided into sections of related disorders, such as disorders of amino acid metabolism, lipid storage disorders and mitochondrial diseases, with an introductory outline where appropriate summarizing the biochemical features and general management issues. Within sections each chapter deals with an individual disease, starting with a useful summary of major phenotypic expression and including clear and helpful biochemical pathways, identifying for the reader exactly where the defect is occurring. Throughout the book, plentiful photographs, often showing extremely rare disorders, are an invaluable aid to diagnosis.

" This text is well organized, and easy to read. The photographs and illustrations are excellent ... the authors have done an exemplary job in producing a text of great value to those for whom it was written - it deserves a place in the library of any institution treating critically ill children." *Annals RCPSC* (April 2000) About the Author William L Nyhan MD, PhD, Professor of Pediatrics, University of California, San Diego, CA, USA; Bruce A Barshop MD, PhD, Professor of Pediatrics, University of California, San Diego, CA, USA; Aida I Al-Aqeel MD, DCH, FRCP, FACMG, Consultant and Head Pediatrics, Medical Genetics and Consultant Endocrinology, Riyadh Military Hospital, Riyadh, Saudi Arabia