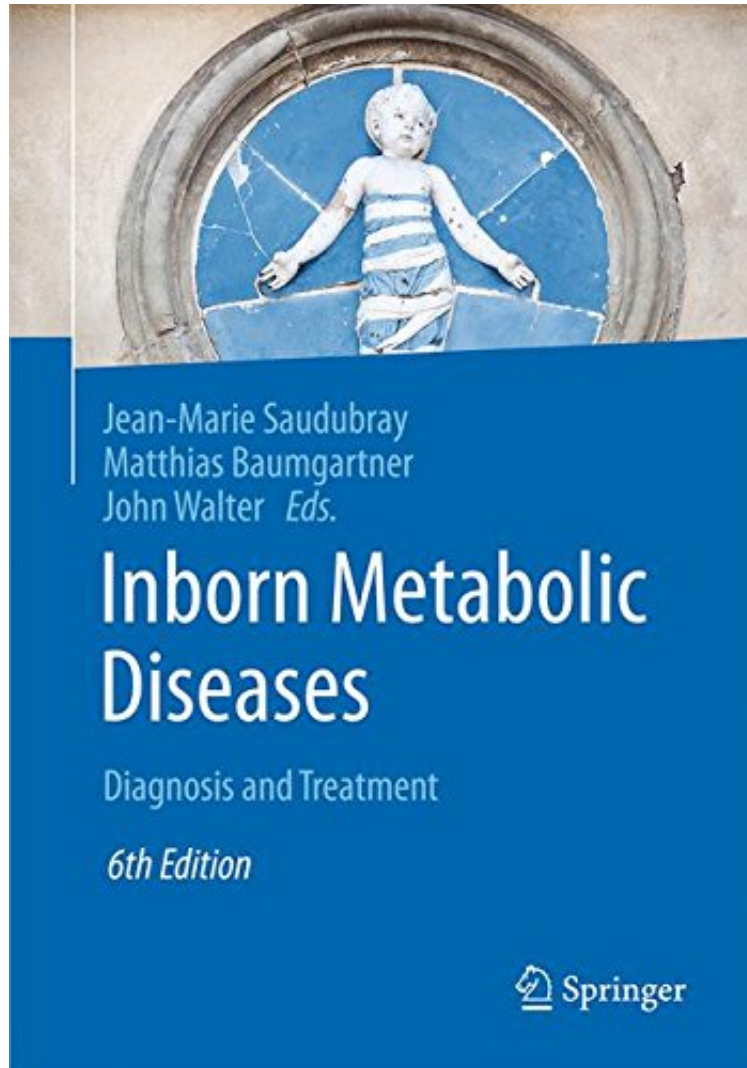


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Inborn Metabolic Diseases: Diagnosis and Treatment remains one of the standard textbooks for professionals working in inborn metabolic medicine and biochemical genetics. Inborn Metabolic Diseases: Diagnosis and Treatment gives an excellent and most up to date overview and deep insight on a rapidly evolving specialty with a steadily growing number of patients, most of them now growing-up and reaching adulthood, and is, therefore, informative for experts as well as for those who seek to become one. (Ulrike Mtze, *Journal of Inherited Metabolic Disease*, Vol. 40, 2017)From the Back CoverThis work is recognised as the standard textbook for professionals involved in the diagnosis and management of inborn errors of metabolism (IEM) and an essential resource in this multidisciplinary field. For the 6th edition all 43 chapters have been newly written or revised by authors with particular expertise in their subject areas. ContentsA clinical and biochemical approach to the recognition and diagnosis of IEM with algorithms to symptoms, signs, and syndromes in patients of all ages; Emergency treatments; Medications. Separate comprehensive sections on IEM of: Carbohydrates; Mitochondrial Energy; Amino and organic acids; Vitamin-responsive defects; Neurotransmitter and Small peptides, Lipid and Bile Acids; Nucleic Acid and Heme; Organelles. Disorders affecting the synthesis and remodelling of complex lipids and fatty acid homeostasis are now included. The Editors Jean-Marie Saudubray, M.D., is Emeritus Professor of Pediatrics and Expert Metabolic Consultant at the adult neuro-metabolic unit in the Hpital La Piti Salptriere from the Universit Pierre et Marie Curie, Paris. Matthias R. Baumgartner, M.D., Professor of Paediatrics and Inherited Metabolic Medicine, University of Zurich, is Head of the Division of Metabolism and the Swiss Newborn Screening Programme at the University Childrens Hospital, Zurich. John Walter, M.D. is Honorary Clinical Professor of Inherited Metabolic Medicine, Manchester Academic Health Science Centre, Willink Unit, Manchester. About the AuthorJean-Marie Saudubray, M.D., is Emeritus Professor of Pediatrics and Expert Metabolic Consultant at the adult neuro-metabolic unit in the Hpital La Piti Salptriere from the Universit Pierre et Marie Curie, Paris. Matthias R. Baumgartner, M.D., Professor of Paediatrics and Inherited Metabolic Medicine, University of Zurich, is Head of the Division of Metabolism and the Swiss Newborn Screening Programme at the University Childrens Hospital, Zurich. John Walter, M.D. is Honorary Clinical Professor of Inherited Metabolic Medicine, Manchester Academic Health Science Centre, Willink Unit, Manchester.