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Altri autori (Persone)	RoycePeter M SteinmannBeat U
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The Marfan Syndrome and Other Microfibrillar Disorders
Chapter 13 The Homocystinurias Chapter 14 Menkes Disease and the Occipital Horn Syndrome; Chapter 15 Epidermolysis Bullosa; Chapter 16 Prolidase Deficiency; Chapter 17 $\alpha(1)$ -Antitrypsin Deficiency; Chapter 18 Heritable Forms of Rickets and Osteomalacia; Chapter 19 Osteopetrosis; Chapter 20 Alkaptonuria; Chapter 21 Fibrodysplasia Ossificans Progressiva; Chapter 22 Disorders of Lysosomal Enzymes; Part I General Considerations; Part II Clinical Phenotypes; Chapter 23 Skeletal Dysplasias; Part I Chondrodysplasias: General Concepts and Diagnostic and Management Considerations
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Sommario/riassunto

The Second Edition of *Connective Tissue and Its Heritable Disorders: Molecular, Genetic, and Medical Aspects* is the definitive reference text in its field, with over 40% more pages on the nature, diagnosis, and treatment of disease than its predecessor. Collecting new research on disorders detailed in the first edition as well as on those previously excluded, editors Peter Royce and Beat Steinmann provide the most up-to-date clinical and scientific information for medical specialists treating affected individuals. Features of this revised and updated volume include detailed reviews of the cl
