Neurocutaneous Disorders - Hemangomas
A Clinical and Diagnostic Approach

Edited by Christos P Panteliadis
University of Thessaloniki, Greece
By (author) Ramis Benjamin
By (author) Hansjorg Cremer
By (author) Christian Hage
By (author) Heymut Omran
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THE BOOK

Neurocutaneous syndromes and hemangomas encompass a substantial number of congenital or hereditary disorders, and present themselves through variable clinical features. Though often complex and multi-systemic, these disorders can mostly be diagnosed by simple visual inspections and strong clinical expertise. The purpose of this book was to compile in a single volume a comprehensive review of the historical perspective, the clinical features, the current knowledge concerning the pathogenesis of each disease, and the diagnostic and therapeutic strategies associated with these challenging disorders. In particular, there is a strong emphasis throughout on the biochemical, molecular, and genetic basis of these syndromes. The international editorial team have drawn upon contributions from colleagues, and from fully referenced information from thousands of articles and abstracts from over 600 publications, thus providing the reader with an outstanding resource for the diagnosis and treatment of neurocutaneous disorders.

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1. Genetics of Neurocutaneous Syndromes
2. Neurocutaneous Disorders (phacomatoses) - Neurobromatosis (Von Recklighausen disease), Tuberous Sclerosis (Bourneville Disease), Encephalofacial Angiomatosis (Sturge-Weber syndrome), Ataxia-Telangiectasia (Louis-Bar syndrome), Angiomatosis of the Retina and the Cerebellum (Von Hippel-Lindau Disease), Linear Naevus Sebaceous, Naevoid Basal Cell Carcinoma (Gorlin-Goltz syndrome), Klippel-Trenaunay Syndrome (Klippel-Trenaunay-Weber syndrome), Incontinentia Pigmenti (Bloch-Sulzberger syndrome) Hypomelanosis of Ito (Incontinentia Pigmenti Achromians), Neurocutaneous Melanosis, Proteus Syndrome, Sjogren-Larsson Syndrome, Ichthyoses, CHILD syndrome, PHACE Syndrome, Refsum Disease (Heredopathis Atactica Polynueuritiformis), Menkes Disease (Kinky Hair Disease, Tricholiodystrophy)

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Tel: +44 (0) 1892 557767 Fax: +44 (0) 1892 530358 Email: info@anshan.co.uk
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