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| Altri autori (Persone) | RivaDaria CuratoloPaolo |
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| Nota di contenuto | <p>""NEUROCUTANEOUS SYNDROMES IN CHILDREN""; ""Contents""; ""Chapter 1 - Embryological basis of the neurocutaneous syndromes""; ""Chapter 2 - Neurocutaneous diseases in children and adolescents: general clinical pattern""; ""Chapter 3 - Hypomelanosis of Ito""; ""Chapter 4 - Vascular malformations and neurocutaneous disorders""; ""Chapter 5 - Neurocutaneous syndromes and hemimegalencephaly""; ""Chapter 6 - Two rare neurocutaneous syndromes: Wyburn-Mason and Proteus""; ""Chapter 7 - Diagnostic criteria and evaluation of patients with tuberous sclerosis complex"" ""Chapter 8 - Epileptic manifestations in tuberous sclerosis"" ""Chapter 9 - Tuberous sclerosis complex: phenotype-genotype correlations""; ""Chapter 10 - Subependymal giant cell astrocytomas and tuberous sclerosis""; ""Chapter 11 - Neurosurgical strategies in the management of subependymal giant cell tumours in tuberous sclerosis complex""; ""Chapter 12 - The neurofibromatoses: clinical manifestations, natural history and management""; ""Chapter 13 - Neurofibromatosis type 1: the dermatological option""; ""Chapter 14 - Clinical molecular genetics of the neurofibromatoses"" ""Chapter 15 - Cognitive-behavioural phenotype and neurobiological basis of neurofibromatosis type 1""""Chapter 16 - Brain tumours in neurofibromatosis type 1""; ""Chapter 17 - Novel therapeutic</p> |

approaches for plexiform neurofibromas in neurofibromatosis type 1"";
""Chapter 18 - Adjuvant therapy for low-grade glioma in
neurofibromatosis type 1""; ""Chapter 19 - Diagnostic and follow-up
protocols for neurofibromatosis types 1 and 2""
