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The second edition of Neurofibromatosis: A Handbook for Patients, Families and Health Care Professionals is much better than a simple handbook. Korf and Rubenstein have expertly woven patient viewpoints and medical concerns related to the combined and distinctive issues of neurofibromatosis (NF) (NF1, NF2, and schwannomatosis) into an outstanding perspective on these genetic conditions.

**Neurofibromatosis: A Handbook for Patients, Families, and**

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**Neurofibromatosis: A Handbook for Patients, Families, and**

Here is the state-of-the-art on recognizing, managing and living with neurofibromatosis (NF) for patients, families, and health care professionals. From new genetic and diagnostic advances, to associated cardiovascular and endocrine abnormalities, to the significant psychosocial impact of NF, the book is packed with clear, practical guidance for understanding and living with this disability.

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Neurofibromatosis 2 (bilateral acoustic neuromas, tumors of cranial nerves and spinal roots, skin manifestations less frequent than in NF1) Multiple café-au-lait spots (an autosomal dominant trait without other features of neurofibromatosis)

## **Handbook of Genetic Counseling/Neurofibromatosis - Type 1**

into Neurofibromatosis type 1 (NF1) and Neurofibromatosis type 2 (NF2), recognizing NF1 as the classic, peripherally-expressed disorder, and NF2 as the intracranial and CNS- affecting disorder. 1 Although NF1 is often described as a rare disease, its rate of 1 in

### **Neurofibromatosis type 1: natural history and impact on**

The actual diagnosis of neurofibromatosis would be made by a medical doctor through the use of findings from an individual's medical history, clinical examination, family history, MRI findings, and/or genetic testing.

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Neurofibromatosis type 1 (NF1) is an autosomal dominant, multisystem disorder affecting approximately 1 in 3500 people. Significant advances in the understanding of the pathophysiology of NF1 have been made in the last decade.

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