



Genetic Instabilities and Hereditary Neurological Diseases

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This book represents the first authoritative review of all neurological diseases related to repeat expansions. Some of the diseases covered in this volume include fragile X syndrome, spino and bulbar muscular atrophy, myotonic dystrophy, spinocerebellar ataxia type 1 and type 7, Huntingtons disease, and Friedreichs ataxia. The book describes investigations into the underlying molecular mechanisms responsible for these syndromes. For students and researchers alike, **Genetic Instabilities and Hereditary Neurological Diseases** serves as a comprehensive treatise covering many aspects of all neurological diseases.

Key Features

- * First authoritative review of neurological diseases related to repeat expansions
- * Description of clinical, human genetics, molecular biological, and biophysical investigations into these syndromes
- * Contributions by most of the principal research teams in the area, edited by world-renowned leaders
- * Lays the background for future investigations on related diseases

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