

Barbro Lindberg

Understanding Rett Syndrome



**A Practical
Guide for
Parents,
Teachers, and
Therapists**



2nd revised edition

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Understanding Rett Syndrome A Practical Guide For Parents Teachers And Therapists

**Carmen María Galvez-
Sánchez, Casandra Isabel
Montoro, Tindara Caprì**



Understanding Rett Syndrome A Practical Guide For Parents Teachers And Therapists:

Understanding Rett Syndrome Barbro Lindberg, 1991 Rett Syndrome is a severe neurological disorder whose cause is presently unknown It leads to a variety of problems often involving mental retardation serious motor handicaps epileptic seizures and difficulties with communication This syndrome is found only in girls and it usually becomes noticeable during their second year of life The disorder is estimated to occur in one out of every 10 000 female births worldwide but as the characteristics of the problem are becoming more widely known new cases are being reported at an increasing rate This is the first book on Rett Syndrome The attempt here is to explain the problem and the implications for the individuals involved their parents teachers and others who work with them Because there is no cure yet medical science presently has little help to offer the families involved although research is proceeding vigorously Many practical insights and techniques are offered for those working with this problem

Rett Syndrome - Clinical and Biological Aspects Bengt Hagberg, Maria Anvret, Jan Wahlström, 1993-07-30 This volume provides a detailed survey of the clinical development of Rett syndrome from its earliest manifestations in childhood to adulthood The volume surveys the developmental profile of the disease its characteristic cluster of symptoms and signs and categorizes the four main clinical stages in the development of motor disability Particular emphasis is given to Rett syndrome variants and other clinical conditions which manifest themselves in a similar way to Rett syndrome The development of scoliosis with age and disease stage is analysed The neurophysiologic neuropathologic and neurochemical characteristics are also examined In addition the volume looks at the molecular genetics of the syndrome drug treatment and possible future developments

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