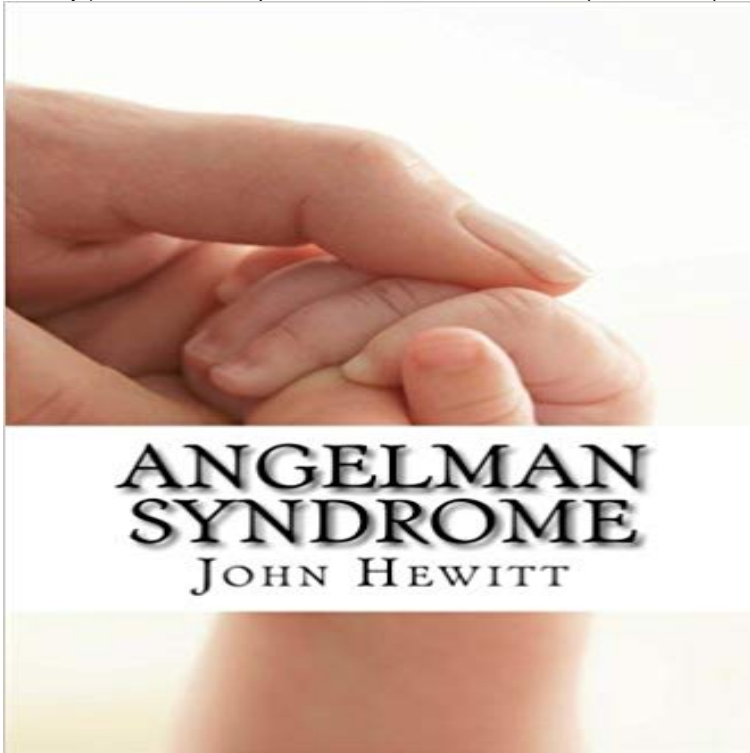


Angelman Syndrome: Causes, Tests, and Treatments



Angelman syndrome is a complex genetic disorder that was first described in 1965 by the English physician Dr. Harry Angelman. At the time, three children who presented a variety of disabilities were admitted to his center. Although at first sight it looked like they were suffering from different disorders, Dr. Angelman felt that there was a common cause for their conditions. He was right. The condition--caused by a chromosomal abnormality--Angelman syndrome is typically diagnosed between the ages of three and seven, and in some cases, it is misdiagnosed for other conditions. This informative book is packed with all you need to know about this devastating condition and includes causes, tests, and treatment options.

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Angelman syndrome is a complex genetic disorder that primarily affects the nervous system. Characteristic features of this condition include **Angelman Syndrome: Causes, Tests and Treatments - Kindle edition** Angelman syndrome is a neurological disorder caused by a missing section of There is no cure, but the child can benefit from treatment, including physical therapy, for the clinical features of Angelman syndrome and performing DNA tests. **Angelman Syndrome: Causes, Tests, and Treatments:** Is there any treatment for Angelman syndrome? What is What are the characteristics, signs, and symptoms of Angelman syndrome? patients and family members the tools to make the right choice in regard to test for a disease or condition. **Angelman Syndrome - GeneReviews - NCBI Bookshelf**

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