

[PDF] Smith's Recognizable Patterns Of Human Malformation: Expert Consult - Online And Print, 7e

Kenneth Lyons Jones, Marilyn C. Jones, Miguel Del Campo - pdf download free book

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1041042 MICRODELETION SYNDROME
(1041) MICRODELETION SYNDROME
Chromosomal Chromosomal Features: Molecular Cytogenetics

Background/History
This syndrome was first described in 1987, associated with the deletion of the short arm of chromosome 5, specifically the region 5p13-5p12. The syndrome is characterized by a variety of clinical features, including growth retardation, facial dysmorphism, and intellectual disability. The syndrome is caused by a deletion of the short arm of chromosome 5, specifically the region 5p13-5p12. The syndrome is characterized by a variety of clinical features, including growth retardation, facial dysmorphism, and intellectual disability. The syndrome is caused by a deletion of the short arm of chromosome 5, specifically the region 5p13-5p12.

ANOMALIES
Growth retardation, facial dysmorphism, intellectual disability, and other clinical features.

Genetics
The syndrome is caused by a deletion of the short arm of chromosome 5, specifically the region 5p13-5p12. The syndrome is characterized by a variety of clinical features, including growth retardation, facial dysmorphism, and intellectual disability. The syndrome is caused by a deletion of the short arm of chromosome 5, specifically the region 5p13-5p12.

OCASIONAL ANOMALIES
Occasional anomalies include growth retardation, facial dysmorphism, and intellectual disability. The syndrome is characterized by a variety of clinical features, including growth retardation, facial dysmorphism, and intellectual disability. The syndrome is caused by a deletion of the short arm of chromosome 5, specifically the region 5p13-5p12.

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Description:

2014 BMA Medical Book Awards 1st Prize Award Winner in Illustrated Book category and Highly Commended in Paediatrics category!

Smith's Recognizable Patterns of Human Malformation has long been known as **the** source to consult on multiple malformation syndromes of environmental and genetic etiology as well as recognizable disorders of unknown cause. This esteemed medical reference **book provides you with complete and authoritative, yet accessible guidance to help accurately diagnose these human disorders, establish prognoses, and provide appropriate management and genetic counseling.**

- Recognize the visual signs of each environmental and genetic abnormality by consulting more than 1,500 full-color photographs and illustrations, many from the personal collections of Drs. Smith and Jones.
 - **Find all the answers you need** about normal and abnormal morphogenesis, minor anomalies and their relevance, clinical approaches to specific diagnoses, and normal standards of measurement for the entire spectrum of human malformation syndromes.
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