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Monosomy 13q Syndrome

National Cancer Institute

Source

National Cancer Institute. *Monosomy 13q Syndrome*. NCI Thesaurus. Code C98993.

A rare syndrome that is characterized by the partial deletion of the long arm of chromosome 13. Signs and symptoms include low birth weight, craniofacial malformations, hands and feet malformations, and mental and psychomotor retardation.