

Fragile X Fragile Hope Finding Joy In Parenting A Child With Special Needs

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## Summary:

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Fragile X syndrome - Wikipedia Fragile X syndrome is typically due to an expansion of the CGG triplet repeat within the Fragile X mental retardation 1 (FMR1) gene on the X chromosome. This results in not enough fragile X mental retardation protein (FMRP), which is required for normal development of the connection between neurons. Fragile X syndrome - Genetics Home Reference - NIH Fragile X syndrome is a genetic condition that causes a range of developmental problems including learning disabilities and cognitive impairment. Usually, males are more severely affected by this disorder than females. What is Fragile X Syndrome? Fragile X Research - FRAXA ... Fragile X syndrome is a rare or orphan disease which affects 1 in 4000 males and 1 in 6000 females. It occurs when a single gene, FMR1, on the long arm of the X chromosome, shuts down and fails to produce a protein, FMRP, which is vital for normal brain development. Fragile X is inherited.

Learning About Fragile X Syndrome - National Human Genome ... Fragile X syndrome is the most common form of inherited intellectual disability in males and is also a significant cause of intellectual disability in females. It affects about 1 in 4,000 males and 1 in 8,000 females and occurs in all racial and ethnic groups. Fragile X Syndrome: Click for Facts, Symptoms, and Treatment Fragile X syndrome (also called Fragile X) is the most common inherited form of mental retardation.\* It results from a change, or mutation, in a single gene, which can be passed from one generation to the next. Fragile X appears in families of every ethnic group and income level. Fragile X Syndrome: Causes, Treatments, and More - WebMD Fragile X syndrome affects a child's learning, behavior, appearance, and health. Symptoms can be mild or more severe. Boys often have a more serious form of it than girls. Children who are born.

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