

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

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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. Fragile X Syndrome: Symptoms, Causes, Diagnosis, and Treatment Boys have one X and one Y chromosome. If the X chromosome has the gene change, they will have symptoms of fragile X syndrome. Some people inherit the fragile X gene without having symptoms. Fragile X syndrome - Wikipedia Fragile X syndrome is a genetic disorder which occurs as a result of a mutation of the fragile X mental retardation 1 (FMR1) gene on the X chromosome, most commonly an increase in the number of CGG trinucleotide repeats in the 5' untranslated region of FMR1.

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: MedlinePlus Fragile X syndrome is the most common form of inherited developmental disability. A problem with a specific gene causes the disease. Normally, the gene makes a protein you need for brain development. What is Fragile X Syndrome? â€¢ Fragile X Research - FRAXA ... A simple DNA blood test for Fragile X syndrome is widely available and very accurate. This test can detect carriers and fully affected individuals. Any doctor can order the test.

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.

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