

Fragile X Syndrome Guide Teachers

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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. Affected individuals usually have delayed development of speech and language by age 2. Fragile X syndrome - Wikipedia Fragile X syndrome (FXS) is a genetic disorder. Symptoms often include mild to moderate intellectual disability. Physical features may include a long and narrow face, large ears, flexible fingers, and large testicles. About. 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.

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. Fragile X Syndrome: Causes, Symptoms, and Diagnosis Fragile X syndrome (FXS) is an inherited genetic disease passed down from parents to children that causes intellectual and developmental disabilities. It's also known as Martin-Bell syndrome. 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.

Facts | Fragile X Syndrome (FXS) | NCBDDD | CDC Fragile X syndrome (FXS) is a genetic disorder. A genetic disorder means that there are changes to the person's genes. FXS is caused by changes in the fragile X mental retardation 1 (FMR1) gene.

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