# Fragile X Syndrome Guide Teachers

**Summary:**



Fragile X syndrome - Genetics Home Reference - NIH Fragile X syndrome is inherited in an X-linked dominant pattern.A condition is considered X-linked if the mutated gene that causes the disorder is located on the X chromosome, one of the two sex chromosomes.(The Y chromosome is the other sex chromosome.) The inheritance is dominant if one copy of the altered gene in each cell is sufficient to cause the condition. Fragile X Syndrome: Symptoms, Causes, Diagnosis, and Treatment Fragile X Syndrome is an inherited disorder caused by genetics that affects a child's learning, behavior, appearance, and health. Learn more about the symptoms, causes, diagnosis, and treatment of. 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 - Wikipedia Fragile X syndrome (FXS) is a genetic disorder. Symptoms often include mild to moderate intellectual disability. The average IQ in males is under 55. Physical features may include a long and narrow face, large ears, flexible fingers, and large testicles. About a third of those affected have features of autism such as problems with social interactions and delayed speech. 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. Itâ€™s also known as Martin-Bell syndrome. Learning About Fragile X Syndrome - genome.gov 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. Home | Fragile X Syndrome (FXS) | NCBDDD | CDC Fragile X syndrome (FXS) is one of the most common causes of inherited intellectual disability. CDC is working to learn more about fragile X syndrome and fragile X-associated disorders to improve the health and well-being of people with these conditions.

fragile x syndrome
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