X-linked hyper IgM syndrome

X-linked hyper IgM syndrome is a condition that affects the immune system and occurs almost exclusively in males. People with this disorder have abnormal levels of proteins called antibodies or immunoglobulins. Antibodies help protect the body against infection by attaching to specific foreign particles and germs, marking them for destruction. There are several classes of antibodies, and each one has a different function in the immune system. Although the name of this condition implies that affected individuals always have high levels of immunoglobulin M (IgM), some people have normal levels of this antibody. People with X-linked hyper IgM syndrome have low levels of three other classes of antibodies: immunoglobulin G (IgG), immunoglobulin A (IgA), and immunoglobulin E (IgE). The lack of certain antibody classes makes it difficult for people with this disorder to fight off infections.

Individuals with X-linked hyper IgM syndrome begin to develop frequent infections in infancy and early childhood. Common infections include pneumonia, sinus infections (sinusitis), and ear infections (otitis). Infections often cause these children to have chronic diarrhea and they fail to gain weight and grow at the expected rate (failure to thrive). Some people with X-linked hyper IgM syndrome have low levels of white blood cells called neutrophils (neutropenia). Affected individuals may develop autoimmune disorders, neurologic complications from brain and spinal cord (central nervous system) infections, liver disease, and gastrointestinal tumors. They also have an increased risk of lymphoma, which is a cancer of immune system cells.

The severity of X-linked hyper IgM syndrome varies among affected individuals, even among members of the same family. Without treatment, this condition can result in death during childhood or adolescence.

Frequency

X-linked hyper IgM syndrome is estimated to occur in 2 per million newborn boys.

Causes

Mutations in the CD40LG gene cause X-linked hyper IgM syndrome. This gene provides instructions for making a protein called CD40 ligand, which is found on the surface of immune system cells known as T cells. CD40 ligand attaches like a key in a lock to its receptor protein, which is located on the surface of immune system cells called B cells. B cells are involved in the production of antibodies, and initially they are able to make only IgM antibodies. When CD40 ligand and its receptor protein are connected, they trigger a series of chemical signals that instruct the B cell to start making IgG, IgA, or IgE antibodies.
CD40 ligand is also necessary for T cells to interact with other cells of the immune system, and it plays a key role in T cell differentiation (the process by which cells mature to carry out specific functions).

Mutations in the CD40LG gene lead to the production of an abnormal CD40 ligand or prevent production of this protein. If CD40 ligand does not attach to its receptor on B cells, these cells cannot produce IgG, IgA, or IgE antibodies. Mutations in the CD40LG gene also impair the T cell's ability to differentiate and interact with other immune system cells. People with X-linked hyper IgM syndrome are more susceptible to infections because they do not have a properly functioning immune system.

**Inheritance Pattern**

This condition is inherited in an X-linked recessive pattern. The gene associated with this condition is located on the X chromosome, which is one of the two sex chromosomes. In males (who have only one X chromosome), one altered copy of the gene in each cell is sufficient to cause the condition. In females (who have two X chromosomes), a mutation would have to occur in both copies of the gene to cause the disorder. Because it is unlikely that females will have two altered copies of this gene, males are affected by X-linked recessive disorders much more frequently than females. A characteristic of X-linked inheritance is that fathers cannot pass X-linked traits to their sons.

**Other Names for This Condition**

- HIGM1
- Hyper-IgM syndrome 1
- Immunodeficiency with Hyper-IgM, type 1

**Diagnosis & Management**

**Genetic Testing Information**

- What is genetic testing? /primer/testing/genetictesting

**Research Studies from ClinicalTrials.gov**

- ClinicalTrials.gov https://clinicaltrials.gov/ct2/results?cond=%22Dysgammaglobulinemia%22+OR+%22X-linked+hyper+IgM+syndrome%22
Other Diagnosis and Management Resources

- GeneReview: X-Linked Hyper IgM Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1402
- MedlinePlus Encyclopedia: Immunodeficiency Disorders
  https://medlineplus.gov/ency/article/000818.htm

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Immunodeficiency Disorders
  https://medlineplus.gov/ency/article/000818.htm
- Health Topic: Immune System and Disorders
  https://medlineplus.gov/immunesystemanddisorders.html

Genetic and Rare Diseases Information Center

- Immunodeficiency with hyper IgM type 1
  https://rarediseases.info.nih.gov/diseases/73/immunodeficiency-with-hyper-igm-type-1

Additional NIH Resources

- National Institute of Allergy and Infectious Diseases: Primary Immune Deficiency Diseases

Educational Resources

- MalaCards: immunodeficiency with hyper-igm, type 1
  https://www.malacards.org/card/immunodeficiency_with_hyper_igm_type_1
- Merck Manual for Healthcare Professionals: Hyper IgM syndrome

Patient Support and Advocacy Resources

- Immune Deficiency Foundation
  https://primaryimmune.org/
- International Patient Organisation for Primary Immunodeficiencies
  https://ipopi.org/
- Jeffrey Modell Foundation: National Primary Immunodeficiency Resource Center
  http://www.info4pi.org/
Clinical Information from GeneReviews
• X-Linked Hyper IgM Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1402

Scientific Articles on PubMed
• PubMed
  +syndrome%5BTIAB%5D%29+OR+%28hyper-igm+syndrome+1%5BTIAB%5D
  %29+OR+%28higm1%5BTIAB%5D%29%29+AND+english%5Bla%5D+AND
  +human%5Bmh%5D+AND+%22last+1440+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM
• IMMUNODEFICIENCY WITH HYPER-IgM, TYPE 1
  http://omim.org/entry/308230

Medical Genetics Database from MedGen
• Immunodeficiency with hyper IgM type 1

Sources for This Summary
• Bhushan A, Covey LR. CD40:CD40L interactions in X-linked and non-X-linked hyper-IgM
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/11817328

• Durandy A. Hyper-IgM syndromes: a model for studying the regulation of class switch
  Review.
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12196205

  Immunological and genetic analysis of 65 patients with a clinical suspicion of X linked hyper-IgM.
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/14514918
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1187335/

• Jain A, Atkinson TP, Lipsky PE, Slater JE, Nelson DL, Strober W. Defects of T-cell effector function
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/10207167
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC408278/

• Johnson J, Filipovich AH, Zhang K. X-Linked Hyper IgM Syndrome. 2007 May 31 [updated 2013
  Jan 24]. In: Pagon RA, Adam MP, Ardinger HH, Wallace SE, Amemiya A, Bean LJH, Bird TD,
  Ledbetter N, Mefford HC, Smith RJH, Stephens K, editors. GeneReviews® [Internet]. Seattle (WA):
  NBK1402/
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20301576

Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/9255191


Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17307885


Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/14663287


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