Autosomal recessive hyper-IgE syndrome

Autosomal recessive hyper-IgE syndrome (AR-HIES) is a disorder of the immune system. A hallmark feature of the condition is recurrent infections that are severe and can be life-threatening. Skin infections can be caused by bacteria, viruses, or fungi. These infections cause rashes, blisters, accumulations of pus (abscesses), open sores, and scaling. People with AR-HIES also tend to have frequent bouts of pneumonia and other respiratory tract infections.

Other immune system-related problems in people with AR-HIES include an inflammatory skin disorder called eczema, food or environmental allergies, and asthma. In some affected individuals, the immune system malfunctions and attacks the body’s own tissues and organs, causing autoimmune disease. For example, autoimmunity can lead to abnormal destruction of red blood cells (hemolytic anemia) in people with AR-HIES.

AR-HIES is characterized by abnormally high levels of an immune system protein called immunoglobulin E (IgE) in the blood; the levels are more than 10 times higher than normal. IgE normally triggers an immune response against foreign invaders in the body, particularly parasitic worms, and plays a role in allergies. It is unclear why people with AR-HIES have such high levels of this protein. People with AR-HIES also have highly elevated numbers of certain white blood cells called eosinophils (hypereosinophilia). Eosinophils aid in the immune response and are involved in allergic reactions.

Some people with AR-HIES have neurological problems, such as paralysis that affects the face or one side of the body (hemiplegia). Blockage of blood flow in the brain or abnormal bleeding in the brain, both of which can lead to stroke, can also occur in AR-HIES.

People with AR-HIES have a greater-than-average risk of developing cancer, particularly cancers of the blood or skin.

Frequency

AR-HIES is a rare disorder whose prevalence is unknown.

Causes

AR-HIES is usually caused by mutations in the DOCK8 gene. The protein produced from this gene plays a critical role in the survival and function of several types of immune system cells. One of the protein’s functions is to help maintain the structure and integrity of immune cells called T cells and NK cells, which recognize and attack foreign invaders, particularly as these cells travel to sites of infection within the body. In addition, DOCK8 is involved in chemical signaling pathways that stimulate other
immune cells called B cells to mature and produce antibodies, which are specialized proteins that attach to foreign particles and germs, marking them for destruction.

DOCK8 gene mutations result in the production of little or no functional DOCK8 protein. Shortage of this protein impairs normal immune cell development and function. It is thought that T cells and NK cells lacking DOCK8 cannot maintain their shape as they move through dense spaces, such as those found within the skin. The abnormal cells die, resulting in reduced numbers of these cells. A shortage of these immune cells impairs the immune response to foreign invaders, accounting for the severe viral skin infections common in AR-HIES. A lack of DOCK8 also impairs B cell maturation and the production of antibodies. A lack of this type of immune response leads to recurrent respiratory tract infections in people with this disorder. It is unclear how DOCK8 gene mutations are involved in other features of AR-HIES, such as the elevation of IgE levels, autoimmunity, and neurological problems.

Some people with AR-HIES do not have mutations in the DOCK8 gene. In these cases, the condition is likely caused by mutations in other genes, some of which have not been identified.

Inheritance Pattern

This condition is inherited in an autosomal recessive pattern, which means both copies of the gene in each cell have mutations. The parents of an individual with an autosomal recessive condition each carry one copy of the mutated gene, but they typically do not show signs and symptoms of the condition.

Other Names for This Condition

- AR-HIES
- autosomal recessive HIES
- CID due to DOCK8 deficiency
- combined immunodeficiency due to DOCK8 deficiency
- DOCK8 deficiency
- DOCK8 immunodeficiency syndrome
- hyper IgE recurrent infection syndrome, autosomal recessive
- hyper immunoglobulin E syndrome, autosomal recessive
- hyperimmunoglobulin E recurrent infection syndrome, autosomal recessive
- hyperimmunoglobulin E syndrome type 2
- non-skeletal hyper-IgE syndrome

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Diagnosis & Management

Genetic Testing Information

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

• Genetic Testing Registry: Hyperimmunoglobulin E syndrome

Research Studies from ClinicalTrials.gov

• ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22autosomal+recessive+hyper-IgE+syndrome%22

Other Diagnosis and Management Resources

• MedlinePlus Encyclopedia: Hyperimmunoglobulin E Syndrome
  https://medlineplus.gov/ency/article/001311.htm

• Merck Manual Professional Version: Hyperimmunoglobulin E Syndrome
  https://www.merckmanuals.com/professional/immunology-allergic-disorders/immunodeficiency-disorders/hyper-ige-syndrome

• PID UK: Hyperimmunoglobulin E Syndromes Treatment and Immunizations
  http://www.piduk.org/specificpidconditions/welldefinedsyndromes/hyperigesyndromes/treatmentandimmunisation

Additional Information & Resources

Health Information from MedlinePlus

• Encyclopedia: Hyperimmunoglobulin E Syndrome
  https://medlineplus.gov/ency/article/001311.htm

• Health Topic: Immune System and Disorders
  https://medlineplus.gov/immunesystemanddisorders.html

• Health Topic: Lung Diseases
  https://medlineplus.gov/lungdiseases.html

• Health Topic: Pneumonia
  https://medlineplus.gov/pneumonia.html

• Health Topic: Skin Infections
  https://medlineplus.gov/skininfections.html

Genetic and Rare Diseases Information Center

• Autosomal recessive hyper IgE syndrome
Additional NIH Resources

- National Institute of Allergy and Infectious Diseases: DOCK8 Deficiency
  https://www.niaid.nih.gov/diseases-conditions/dock8-deficiency
- National Institute of Allergy and Infectious Diseases: Primary Immune Deficiency Diseases

Educational Resources

- MalaCards: hyper-ige recurrent infection syndrome, autosomal recessive
  https://www.malacards.org/card/hyper_ige_recurrent_infection_syndrome_autosomal_recessive
- Orphanet: Combined immunodeficiency due to DOCK8 deficiency
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=217390
- Orphanet: OBSOLETE: Autosomal recessive hyper-IgE syndrome
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=169446

Patient Support and Advocacy Resources

- Immune Deficiency Foundation
  https://primaryimmune.org/about-primary-immunodeficiencies/specific-disease-types/hyper-ige-syndrome/
- International Patient Organisation for Primary Immunodeficiencies
  https://ipopi.org/
- Jeffrey Modell Foundation
  http://www.info4pi.org/
- National Organization for Rare Disorders
  https://rarediseases.org/rare-diseases/autosomal-recessive-hyper-ige-syndrome/
- PID UK: Hyperimmunoglobulin E Syndromes
  http://www.piduk.org/specificpidconditions/welldefinedsyndromes/hyperigesyndromes
- Resource List from the University of Kansas Medical Center: Immune Deficiency Conditions
  http://www.kumc.edu/gec/support/immune.html
Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28autosomal+recessive+hyper-IgE+syndrome%5BTIAB%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- HYPER-IgE RECURRENT INFECTION SYNDROME 2, AUTOSOMAL RECESSIVE
  http://omim.org/entry/243700

Sources for This Summary


  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/29907690
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6173313/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/22581261
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  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4016982/

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