Adenosine deaminase deficiency

Adenosine deaminase (ADA) deficiency is an inherited disorder that damages the immune system and causes severe combined immunodeficiency (SCID). People with SCID lack virtually all immune protection from bacteria, viruses, and fungi. They are prone to repeated and persistent infections that can be very serious or life-threatening. These infections are often caused by "opportunistic" organisms that ordinarily do not cause illness in people with a normal immune system.

The main symptoms of ADA deficiency are pneumonia, chronic diarrhea, and widespread skin rashes. Affected children also grow much more slowly than healthy children and some have developmental delay.

Most individuals with ADA deficiency are diagnosed with SCID in the first 6 months of life. Without treatment, these babies usually do not survive past age 2. In about 10 percent to 15 percent of cases, onset of immune deficiency is delayed to between 6 and 24 months of age (delayed onset) or even until adulthood (late onset). Immune deficiency in these later-onset cases tends to be less severe, causing primarily recurrent upper respiratory and ear infections. Over time, affected individuals may develop chronic lung damage, malnutrition, and other health problems.

Frequency

Adenosine deaminase deficiency is very rare and is estimated to occur in approximately 1 in 200,000 to 1,000,000 newborns worldwide. This disorder is responsible for approximately 15 percent of SCID cases.

Causes

Adenosine deaminase deficiency is caused by mutations in the ADA gene. This gene provides instructions for producing the enzyme adenosine deaminase. This enzyme is found throughout the body but is most active in specialized white blood cells called lymphocytes. These cells protect the body against potentially harmful invaders, such as bacteria and viruses, by making immune proteins called antibodies or by directly attacking infected cells. Lymphocytes are produced in specialized lymphoid tissues including the thymus, which is a gland located behind the breastbone, and lymph nodes, which are found throughout the body. Lymphocytes in the blood and in lymphoid tissues make up the immune system.

The function of the adenosine deaminase enzyme is to eliminate a molecule called deoxyadenosine, which is generated when DNA is broken down. Adenosine deaminase converts deoxyadenosine, which can be toxic to lymphocytes, to another molecule called deoxyinosine that is not harmful. Mutations in the ADA gene reduce or eliminate
the activity of adenosine deaminase and allow the buildup of deoxyadenosine to levels that are toxic to lymphocytes.

Immature lymphocytes in the thymus are particularly vulnerable to a toxic buildup of deoxyadenosine. These cells die before they can mature to help fight infection. The number of lymphocytes in other lymphoid tissues is also greatly reduced. The loss of infection-fighting cells results in the signs and symptoms of SCID.

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
• ADA deficiency
• ADA-SCID
• adenosine deaminase deficient severe combined immunodeficiency
• SCID due to ADA deficiency
• severe combined immunodeficiency due to ADA deficiency
• severe combined immunodeficiency, autosomal recessive, T cell-negative, B cell-negative, NK cell-negative, due to adenosine deaminase deficiency

Diagnosis & Management
Formal Diagnostic Criteria
• ACT Sheet: Severe Combined Immunodeficiency (SCID) and Conditions Associated with T Cell Lymphoeia
  https://www.ncbi.nlm.nih.gov/books/NBK55827/bin/SCID.pdf

Genetic Testing Information
• What is genetic testing?
  /primer/testing/genetictesting
• Genetic Testing Registry: Severe combined immunodeficiency due to ADA deficiency

Research Studies from ClinicalTrials.gov
• ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22Severe+Combined+Immunodeficiency%22+OR+%22adenosine+deaminase+deficiency%22
Other Diagnosis and Management Resources

- American Society of Gene and Cell Therapy: Inherited Immunodeficiencies
  https://www.asgct.org/education/disease-treatments/inherited-immunodeficiencies

- Baby’s First Test: Severe Combined Immunodeficiency
  https://www.babysfirsttest.org/newborn-screening/conditions/severe-combined-immunodeficiency-scid

- GeneReview: Adenosine Deaminase Deficiency
  https://www.ncbi.nlm.nih.gov/books/NBK1483

- National Marrow Donor Program: SCID and Transplant

Additional Information & Resources

Health Information from MedlinePlus

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

- Health Topic: Newborn Screening
  https://medlineplus.gov/newbornscreening.html

Genetic and Rare Diseases Information Center

- Adenosine deaminase deficiency
  https://rarediseases.info.nih.gov/diseases/5748/adenosine-deaminase-deficiency

Additional NIH Resources

- National Human Genome Research Institute: Learning About Severe Combined Immunodeficiency
  https://www.genome.gov/13014325/

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

Educational Resources

- KidsHealth

- MalaCards: adenosine deaminase deficiency
  https://www.malacards.org/card/adenosine_deaminase_deficiency

- Merck Manual Consumer Version
  https://www.merckmanuals.com/home/immune-disorders/immunodeficiency-disorders/severe-combined-immunodeficiency-scid
- Orphanet: Severe combined immunodeficiency due to adenosine deaminase deficiency
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=277
- University of Utah Genetic Science Learning Center
  https://learn.genetics.utah.edu/content/disorders/singlegene/

Patient Support and Advocacy Resources
- Children Living with Inherited Metabolic Diseases
  https://www.metabolicsupportuk.org/
- Immune Deficiency Foundation
  https://primaryimmune.org/
- Jeffrey Modell Foundation
  http://www.info4pi.org/
- National Organization for Rare Disorders (NORD)
  https://rarediseases.org/rare-diseases/severe-combined-immunodeficiency/

Clinical Information from GeneReviews
- Adenosine Deaminase Deficiency
  https://www.ncbi.nlm.nih.gov/books/NBK1483

Scientific Articles on PubMed
- PubMed
deficiency%5BMAJR%5D%29+AND+%28%28adenosine+deaminase+deficiency%5BMAJR%5D%29+OR+%28%28ada-scid%5BMAJR%5D%29+OR+%28%28ade
mosine+deaminase+deficient+severe+combined+immunodeficiency%5BMAJR%5D%29+OR+%28%28ada+deficiency%5BMAJR%5D%29%29+AND+english%5BMAJR%5D+AND
+human%5BMAJR%5D+AND+%22last+1800+days%22+AND+pubmedterm+AND+ad

Catalog of Genes and Diseases from OMIM
- SEVERE COMBINED IMMUNODEFICIENCY, AUTOSOMAL RECESSIVE, T CELL-NEGATIVE, B CELL-NEGATIVE, NK CELL-NEGATIVE, DUE TO
  ADENOSINE DEAMINASE DEFICIENCY
  http://omim.org/entry/102700
Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/22262755
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3341658/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15032591

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/14499267

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15580654

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17765813

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16176880

Reprinted from Genetics Home Reference:

Reviewed: July 2013
Published: October 9, 2018

Lister Hill National Center for Biomedical Communications
U.S. National Library of Medicine
National Institutes of Health
Department of Health & Human Services