Monoamine oxidase A deficiency

Monoamine oxidase A deficiency is a rare disorder that occurs almost exclusively in males. It is characterized by mild intellectual disability and behavioral problems beginning in early childhood.

Most boys with monoamine oxidase A deficiency are less able to control their impulses than their peers, causing aggressive or violent outbursts. In addition, affected individuals may have features of other behavioral disorders, including autism spectrum disorder and attention-deficit/hyperactivity disorder (ADHD). These features can include obsessive behaviors, difficulty forming friendships, and problems focusing attention. Sleep problems, such as trouble falling asleep or night terrors, can also occur in monoamine oxidase A deficiency.

Some people with monoamine oxidase A deficiency have episodes of skin flushing, sweating, headaches, or diarrhea. Similar episodes can occur in female family members of males with monoamine oxidase A deficiency, although females do not experience other signs or symptoms of the condition.

In some cases, certain foods, such as cheese, appear to worsen symptoms of monoamine oxidase A deficiency.

Frequency

Monoamine oxidase A deficiency is thought to be very rare. Its prevalence is unknown.

Causes

Monoamine oxidase A deficiency is caused by mutations in the MAOA gene. This gene provides instructions for making an enzyme called monoamine oxidase A. This enzyme breaks down chemicals called monoamines, including serotonin, epinephrine, and norepinephrine. These particular monoamines act as neurotransmitters, which transmit signals between nerve cells in the brain. Monoamine oxidase A helps break down the neurotransmitters when signaling is no longer needed. Signals transmitted by serotonin regulate mood, emotion, sleep, and appetite. Epinephrine and norepinephrine control the body’s response to stress. Monoamine oxidase A also helps break down monoamines found in the diet.

Mutations in the MAOA gene reduce monoamine oxidase A activity, which causes serotonin and other neurotransmitters to build up in the brain. It is unclear how this buildup leads to the signs and symptoms of monoamine oxidase A deficiency. Researchers suspect that high levels of serotonin may impair an affected individual's ability to control his impulses, leading to aggressive outbursts. In addition, the
outbursts may be an overreaction to stress, possibly due to the impaired breakdown of epinephrine and norepinephrine.

A reduction of monoamine oxidase A activity also impairs breakdown of monoamines found in foods. An excess of these molecules can contribute to the behavioral problems, flushing, sweating, and other symptoms associated with monoamine oxidase A deficiency, which may be why foods high in monoamines sometimes worsen the symptoms of the condition.

Monoamine oxidase A plays a role in normal brain development. Some studies suggest that reduced monoamine oxidase A activity alters the development of certain regions of the brain, which may contribute to intellectual disability and behavioral problems in people with monoamine oxidase A deficiency.

Research suggests that environmental factors, such as mistreatment in childhood, may impact the severity of the condition and the behavioral problems that develop.

**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**

- Brunner syndrome
- deficiency of monoamine oxidase A
- X-linked monoamine oxidase deficiency

**Diagnosis & Management**

**Genetic Testing Information**

- What is genetic testing? /primer/testing/genetictesting
Additional Information & Resources

Health Information from MedlinePlus
- Encyclopedia: Intellectual Disability
  https://medlineplus.gov/ency/article/001523.htm
- Health Topic: Attention Deficit Hyperactivity Disorder
  https://medlineplus.gov/attentiondeficithyperactivitydisorder.html
- Health Topic: Autism Spectrum Disorder
  https://medlineplus.gov/autismspectrumdisorder.html
- Health Topic: Developmental Disabilities
  https://medlineplus.gov/developmentaldisabilities.html

Genetic and Rare Diseases Information Center
- Monoamine oxidase A deficiency
  https://rarediseases.info.nih.gov/diseases/3531/monoamine-oxidase-a-deficiency

Additional NIH Resources
- MentalHealth.gov: Behavioral Disorders
  https://www.mentalhealth.gov/what-to-look-for/behavioral-disorders
- National Institute of Neurologic Disorders and Stroke: Autism Spectrum Disorder Information Page
  https://www.ninds.nih.gov/Disorders/All-Disorders/Autism-Spectrum-Disorder-Information-Page

Educational Resources
- Centers For Disease Control and Prevention: Facts About Intellectual Disability
- MalaCards: brunner syndrome
  https://www.malacards.org/card/brunner_syndrome
- Orphanet: Monoamine oxidase A deficiency
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=3057
Patient Support and Advocacy Resources

- American Association on Intellectual and Developmental Disabilities (AAIDD)
  https://www.aaidd.org/
- Autism Society
  https://www.autism-society.org/
- Autism Speaks
  https://www.autismspeaks.org/

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28monoamine+oxidase+A+deficiency%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1440+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- BRUNNER SYNDROME
  http://omim.org/entry/300615

Medical Genetics Database from MedGen

- Monoamine oxidase A deficiency

Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/21971001
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3371272/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/8211186

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/7792602
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2844866/
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/25807999

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/24169519
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4023218/

Reprinted from Genetics Home Reference:

Reviewed: May 2017
Published: November 12, 2019

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