Rett syndrome

Rett syndrome is a brain disorder that occurs almost exclusively in girls. The most common form of the condition is known as classic Rett syndrome. After birth, girls with classic Rett syndrome have 6 to 18 months of apparently normal development before developing severe problems with language and communication, learning, coordination, and other brain functions. Early in childhood, affected girls lose purposeful use of their hands and begin making repeated hand wringing, washing, or clapping motions. They tend to grow more slowly than other children and about three-quarters have a small head size (microcephaly). Other signs and symptoms that can develop include breathing abnormalities, spitting or drooling, unusual eye movements such as intense staring or excessive blinking, cold hands and feet, irritability, sleep disturbances, seizures, and an abnormal side-to-side curvature of the spine (scoliosis).

Researchers have described several variant or atypical forms of Rett syndrome, which can be milder or more severe than the classic form.

Rett syndrome is part of a spectrum of disorders with the same genetic cause. Other disorders on the spectrum include PPM-X syndrome, MECP2 duplication syndrome, and MECP2-related severe neonatal encephalopathy. These other conditions can affect males.

Frequency

This condition affects an estimated 1 in 9,000 to 10,000 females.

Causes

Mutations in a gene called MECP2 underlie almost all cases of classic Rett syndrome and some variant forms of the condition. This gene provides instructions for making a protein (MeCP2) that is critical for normal brain function. Although the exact function of the MeCP2 protein is unclear, it is likely involved in maintaining connections (synapses) between nerve cells (neurons). It may also be necessary for the normal function of other types of brain cells.

The MeCP2 protein is thought to help regulate the activity of genes in the brain. This protein may also control the production of different versions of certain proteins in brain cells. Mutations in the MECP2 gene alter the MeCP2 protein or result in the production of less protein, which appears to disrupt the normal function of neurons and other cells in the brain. Specifically, studies suggest that changes in the MeCP2 protein may reduce the activity of certain neurons and impair their ability to communicate with one another. It is unclear how these changes lead to the specific features of Rett syndrome.
Several conditions with signs and symptoms overlapping those of Rett syndrome have been found to result from mutations in other genes. These conditions, including FOXG1 syndrome and CDKL5 deficiency disorder, were previously thought to be variant forms of Rett syndrome. However, doctors and researchers have identified some important differences between the conditions, so they are now usually considered to be separate disorders.

**Inheritance Pattern**

In more than 99 percent of people with Rett syndrome, there is no history of the disorder in their family. Many of these cases result from new mutations in the MECP2 gene.

A few families with more than one affected family member have been described. These cases helped researchers determine that classic Rett syndrome and variants caused by MECP2 gene mutations have an X-linked dominant pattern of inheritance. 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 inheritance is dominant if one copy of the altered gene in each cell is sufficient to cause the condition.

Males with mutations in the MECP2 gene often die in infancy. However, a small number of males with a genetic change involving MECP2 have developed signs and symptoms similar to those of Rett syndrome, including intellectual disability, seizures, and movement problems. In males, this condition is described as MECP2-related severe neonatal encephalopathy. The signs and symptoms in some males with an MECP2 gene mutation are on the milder end of the spectrum.

**Other Names for This Condition**

- autism-dementia-ataxia-loss of purposeful hand use syndrome
- Rett disorder
- Rett's disorder
- Rett's syndrome
- RTT

**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=%22Rett+syndrome%22

Other Diagnosis and Management Resources

- Boston Children's Hospital
  http://www.childrenshospital.org/conditions-and-treatments/conditions/r/rett-syndrome
- GeneReview: MECP2-Related Disorders
  https://www.ncbi.nlm.nih.gov/books/NBK1497
- MedlinePlus Encyclopedia: Rett Syndrome
  https://medlineplus.gov/ency/article/001536.htm
- RettSyndrome.org: Rett Syndrome Clinics
  https://www.rettsyndrome.org/about-rett-syndrome/clinics

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Rett Syndrome
  https://medlineplus.gov/ency/article/001536.htm
- Health Topic: Rett Syndrome
  https://medlineplus.gov/rettsyndrome.html

Genetic and Rare Diseases Information Center

- Atypical Rett syndrome
- Rett syndrome

Additional NIH Resources

- Eunice Kennedy Shriver National Institute of Child Health & Human Development
  https://www.nichd.nih.gov/health/topics/rett
- National Institute of Neurological Disorders and Stroke
  https://www.ninds.nih.gov/Disorders/All-Disorders/Rett-Syndrome-Information-Page

Educational Resources

- InterRett: International Rett Syndrome Database
- Kennedy Krieger Institute
  https://www.kennedykrieger.org/patient-care/conditions/rett-syndrome
• MalaCards: rett syndrome
  https://www.malacards.org/card/rett Syndrome

• Orphanet: Rett syndrome
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=778

• Swedish Information Center for Rare Diseases
  http://www.socialstyrelsen.se/rarediseases/rettsyndrome

Patient Support and Advocacy Resources

• National Organization for Rare Disorders
  https://rarediseases.org/rare-diseases/rett-syndrome/

• RareConnect

• Resource List from the University of Kansas Medical Center
  http://www.kumc.edu/gec/support/rett_syn.html

• Rett Syndrome Association UK
  http://www.rettuk.org/

• Rett Syndrome Research Trust
  https://reverserett.org/

• RettSyndrome.org
  https://www.rettsyndrome.org/

Clinical Information from GeneReviews

• MECP2-Related Disorders
  https://www.ncbi.nlm.nih.gov/books/NBK1497

Scientific Articles on PubMed

• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28Rett+Syndrome%5BMAJR%5D %29+AND+%28Rett+syndrome%5BTL%5D%29+AND+english%5Bla%5D+AND +human%5Bmh%5D+AND+%22last+360+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

• RETT SYNDROME
  http://omim.org/entry/312750
Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17988628

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20301670

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/27884167 
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5123333/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/29185709

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/21154482 
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3058521/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15070486

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16225824

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/21632916 
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3127460/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19238181

Reprinted from Genetics Home Reference: 

Reviewed: October 2018
Published: March 19, 2019