Tetrasomy 18p

Tetrasomy 18p is a chromosomal condition that affects many parts of the body. This condition usually causes feeding difficulties in infancy, delayed development, intellectual disability that is often mild to moderate but can be severe, changes in muscle tone, distinctive facial features, and other birth defects. However, the signs and symptoms vary among affected individuals.

Babies with tetrasomy 18p often have trouble feeding and may vomit frequently, which makes it difficult for them to gain weight. Some affected infants also have breathing problems and jaundice, which is a yellowing of the skin and the whites of the eyes.

Changes in muscle tone are commonly seen with tetrasomy 18p. Some affected children have weak muscle tone (hypotonia), while others have increased muscle tone (hypertonia) and stiffness (spasticity). These changes contribute to delayed development of motor skills, including sitting, crawling, and walking.

Tetrasomy 18p is associated with a distinctive facial appearance that can include unusually shaped and low-set ears, a small mouth, a flat area between the upper lip and the nose (philtrum), and a thin upper lip. Many affected individuals also have a high, arched roof of the mouth (palate), and a few have had a split in the roof of the mouth (cleft palate).

Additional features of tetrasomy 18p can include seizures, vision problems, recurrent ear infections, mild to moderate hearing loss, constipation and other gastrointestinal problems, abnormal curvature of the spine (scoliosis or kyphosis), a shortage of growth hormone, and birth defects affecting the heart and other organs. Males with tetrasomy 18p may be born with undescended testes (cryptorchidism) or the opening of the urethra on the underside of the penis (hypospadias). Psychiatric conditions, such as attention-deficit/hyperactivity disorder (ADHD) and anxiety, as well as social and behavioral challenges have also been reported in some people with tetrasomy 18p.

Frequency

Tetrasomy 18p is a rare disorder. It is known to affect about 250 families worldwide.

Causes

Tetrasomy 18p results from the presence of an abnormal extra chromosome, called an isochromosome 18p, in each cell. An isochromosome is a chromosome with two identical arms. Normal chromosomes have one long (q) arm and one short (p) arm, but isochromosomes have either two q arms or two p arms. Isochromosome 18p is a version of chromosome 18 made up of two p arms.
Cells normally have two copies of each chromosome, one inherited from each parent.
In people with tetrasomy 18p, cells have the usual two copies of chromosome 18
plus an isochromosome 18p. As a result, each cell has four copies of the short arm
of chromosome 18. (The word "tetrasomy" is derived from "tetra," the Greek word for
"four.") The extra genetic material from the isochromosome disrupts the normal course
of development, causing the characteristic features of this disorder.

**Inheritance Pattern**

Tetrasomy 18p is usually not inherited. The chromosomal change responsible for the
disorder typically occurs as a random event during the formation of reproductive cells
(eggs or sperm) in a parent of the affected individual, usually the mother. Most affected
individuals have no history of the disorder in their family. However, rare inherited cases
of tetrasomy 18p have been reported.

**Other Names for This Condition**

- 18p isochromosome
- 18p tetrasomy

**Diagnosis & Management**

**Genetic Testing Information**

- What is genetic testing? /primer/testing/genetictesting
- Genetic Testing Registry: Chromosome 18, tetrasomy 18p

**Other Diagnosis and Management Resources**

- Chromosome 18 Clinical Research Center, University of Texas Health Science
  Center at San Antonio
  https://wp.uthscsa.edu/chrome-18/

**Additional Information & Resources**

**Health Information from MedlinePlus**

- Encyclopedia: Chromosome
  https://medlineplus.gov/ency/article/002327.htm
- Health Topic: Developmental Disabilities
  https://medlineplus.gov/developmentaldisabilities.html

**Genetic and Rare Diseases Information Center**

- Chromosome 18p tetrasomy
  https://rarediseases.info.nih.gov/diseases/35/chromosome-18p-tetrasomy
Educational Resources

- MalaCards: chromosome 18p tetrasomy
  https://www.malacards.org/card/chromosome_18p_tetrasomy

- March of Dimes: Chromosomal Conditions
  https://www.marchofdimes.org/baby/chromosomal-conditions.aspx

- Orphanet: Tetrasomy 18p
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=3307

Patient Support and Advocacy Resources

- Chromosome 18 Registry & Research Society
  https://www.chromosome18.org/tetrasomy-18p/

- Chromosome Disorder Outreach
  https://chromodisorder.org/

- National Organization for Rare Disorders (NORD)
  https://rarediseases.org/rare-diseases/chromosome-18-tetrasomy-18p/

- Resource list from the University of Kansas Medical Center: Chromosomal Conditions
  http://www.kumc.edu/gec/support/chromoso.html

- Unique - The Rare Chromosome Disorder Support Group (UK)
  https://www.rarechromo.org/

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28tetrasomy+18p%5BTIAB%5D%29+OR+%28isochromosome+18p%5BTIAB%5D%29+AND+english%5BLanguage%5D+AND+human%5BMedical_Human%5D+AND+%22last+3600+days%22%5BDate%5D

Catalog of Genes and Diseases from OMIM

- TETRASOMY 18p
  http://omim.org/entry/614290

Sources for This Summary


  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/25900901

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20803640

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/25403900

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19927307


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