Familial restrictive cardiomyopathy

Familial restrictive cardiomyopathy is a genetic form of heart disease. For the heart to beat normally, the heart (cardiac) muscle must contract and relax in a coordinated way. Oxygen-rich blood from the lungs travels first through the upper chambers of the heart (the atria), and then to the lower chambers of the heart (the ventricles).

In people with familial restrictive cardiomyopathy, the heart muscle is stiff and cannot fully relax after each contraction. Impaired muscle relaxation causes blood to back up in the atria and lungs, which reduces the amount of blood in the ventricles.

Familial restrictive cardiomyopathy can appear anytime from childhood to adulthood. The first signs and symptoms of this condition in children are failure to gain weight and grow at the expected rate (failure to thrive), extreme tiredness (fatigue), and fainting. Children who are severely affected may also have abnormal swelling or puffiness (edema), increased blood pressure, an enlarged liver, an abnormal buildup of fluid in the abdominal cavity (ascites), and lung congestion. Some children with familial restrictive cardiomyopathy do not have any obvious signs or symptoms, but they may die suddenly due to heart failure. Without treatment, the majority of affected children survive only a few years after they are diagnosed.

Adults with familial restrictive cardiomyopathy typically first develop shortness of breath, fatigue, and a reduced ability to exercise. Some individuals have an irregular heart beat (arrhythmia) and may also experience a sensation of fluttering or pounding in the chest (palpitations) and dizziness. Abnormal blood clots are commonly seen in adults with this condition. Without treatment, approximately one-third of adults with familial restrictive cardiomyopathy do not survive more than five years after diagnosis.

Frequency

The prevalence of familial restrictive cardiomyopathy is unknown. Although cardiomyopathy is a relatively common condition, restrictive cardiomyopathy, in which relaxation of the heart muscle is impaired, is the least common type. Some other forms of cardiomyopathy involve a weak or enlarged heart muscle with impaired contraction. In the United States and in Europe, restrictive cardiomyopathy accounts for less than five percent of all cardiomyopathies. The proportion of restrictive cardiomyopathy that runs in families is not known.

Causes

Mutations in several genes have been found to cause familial restrictive cardiomyopathy. Mutations in the TNNI3 gene are one of the major causes of this condition. The TNNI3 gene provides instructions for making a protein called cardiac troponin I, which is found solely in the heart. Cardiac troponin I is one of three proteins
that make up the troponin protein complex, which helps regulate tensing (contraction) and relaxation of the heart muscle.

*TNNI3* gene mutations associated with familial restrictive cardiomyopathy result in the production of a defective cardiac troponin I protein. The altered protein disrupts the function of the troponin protein complex and does not allow the heart muscle to fully relax. As a result, not enough blood enters the ventricles, leading to a buildup in the atria and lungs. The abnormal heart relaxation and blood flow is responsible for many of the signs and symptoms of familial restrictive cardiomyopathy.

Mutations in other genes associated with familial restrictive cardiomyopathy each account for a small percentage of cases of this condition. Some people with familial restrictive cardiomyopathy do not have an identified mutation in any of the known associated genes. The cause of the disorder in these individuals is unknown.

**Inheritance Pattern**

This condition is inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to cause the disorder.

**Other Names for This Condition**

- cardiomyopathy, restrictive
- RCM

**Diagnosis & Management**

**Genetic Testing Information**

- What is genetic testing?
  /primer/testing/genetictesting
- Genetic Testing Registry: Familial restrictive cardiomyopathy
  https://www.ncbi.nlm.nih.gov/gtr/conditions/C0340429/
- Genetic Testing Registry: Familial restrictive cardiomyopathy 1
- Genetic Testing Registry: Familial restrictive cardiomyopathy 2
- Genetic Testing Registry: Familial restrictive cardiomyopathy 3

**Research Studies from ClinicalTrials.gov**

- ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22familial+restrictive+cardiomyopathy%22+OR+%22Cardiomyopathy%22+OR+%22Restrictive%22
Other Diagnosis and Management Resources

- Johns Hopkins Medicine: Cardiomyopathy
  https://www.hopkinsmedicine.org/healthlibrary/conditions/adult/cardiovascular_diseases/cardiomyopathy_85,p00201
- MedlinePlus Encyclopedia: Restrictive Cardiomyopathy
  https://medlineplus.gov/ency/article/000189.htm

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Restrictive Cardiomyopathy
  https://medlineplus.gov/ency/article/000189.htm
- Health Topic: Cardiomyopathy
  https://medlineplus.gov/cardiomyopathy.html

Additional NIH Resources

- National Heart Lung and Blood Institute: What Is Cardiomyopathy?
  https://www.nhlbi.nih.gov/health-topics/cardiomyopathy

Educational Resources

- American Heart Association: Restrictive Cardiomyopathy
- Boston Children's Hospital: Cardiomyopathy
  http://www.childrenshospital.org/conditions-and-treatments/conditions/c/cardiomyopathy
- Centers for Disease Control and Prevention: Heart Disease
  https://www.cdc.gov/heartdisease/
- Cincinnati Children's Hospital: Cardiomyopathies
  https://www.cincinnatichildrens.org/health/c/cardiomyopathy
- Merck Manual Home Edition for Patients and Caregivers
- Orphanet: Familial restrictive cardiomyopathy
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=217635
Patient Support and Advocacy Resources

- **American Heart Association**
  https://www.heart.org/

- **Children's Cardiomyopathy Foundation**
  http://www.childrenscardiomyopathy.org/

- **National Organization for Rare Disorders (NORD): Pediatric Cardiomyopathy**
  https://rarediseases.org/rare-diseases/pediatric-cardiomyopathy/

- **University of Kansas Medical Center Resource List: Heart / Cardiology Conditions**
  http://www.kumc.edu/gec/support/conghart.html

Scientific Articles on PubMed

- **PubMed**
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28Cardiomyopathy%2C+Restrictive%5BMAJR%5D%29+AND+%28restrictive+cardiomyopathy%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

- **CARDIOMYOPATHY, FAMILIAL RESTRICTIVE, 1**
  http://omim.org/entry/115210

- **CARDIOMYOPATHY, FAMILIAL RESTRICTIVE, 2**
  http://omim.org/entry/609578

- **CARDIOMYOPATHY, FAMILIAL RESTRICTIVE, 3**
  http://omim.org/entry/612422

Sources for This Summary

  *Citation on PubMed:* https://www.ncbi.nlm.nih.gov/pubmed/17916581

  *Citation on PubMed:* https://www.ncbi.nlm.nih.gov/pubmed/18467357

- **Mogensen J, Arbustini E. Restrictive cardiomyopathy.**
  *Citation on PubMed:* https://www.ncbi.nlm.nih.gov/pubmed/19593902

- **Nihoyannopoulos P, Dawson D. Restrictive cardiomyopathies.**
  *Citation on PubMed:* https://www.ncbi.nlm.nih.gov/pubmed/19889655
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20347786

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20298698


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