Vitelliform macular dystrophy

Vitelliform macular dystrophy is a genetic eye disorder that can cause progressive vision loss. This disorder affects the retina, the specialized light-sensitive tissue that lines the back of the eye. Specifically, vitelliform macular dystrophy disrupts cells in a small area near the center of the retina called the macula. The macula is responsible for sharp central vision, which is needed for detailed tasks such as reading, driving, and recognizing faces.

Vitelliform macular dystrophy causes a fatty yellow pigment (lipofuscin) to build up in cells underlying the macula. Over time, the abnormal accumulation of this substance can damage cells that are critical for clear central vision. As a result, people with this disorder often lose their central vision, and their eyesight may become blurry or distorted. Vitelliform macular dystrophy typically does not affect side (peripheral) vision or the ability to see at night.

Researchers have described two forms of vitelliform macular dystrophy with similar features. The early-onset form (known as Best disease) usually appears in childhood; the onset of symptoms and the severity of vision loss vary widely. The adult-onset form begins later, usually in mid-adulthood, and tends to cause vision loss that worsens slowly over time. The two forms of vitelliform macular dystrophy each have characteristic changes in the macula that can be detected during an eye examination.

Frequency
Vitelliform macular dystrophy is a rare disorder; its incidence is unknown.

Causes
Mutations in the BEST1 and PRPH2 genes cause vitelliform macular dystrophy. BEST1 mutations are responsible for Best disease and for some cases of the adult-onset form of vitelliform macular dystrophy. Changes in the PRPH2 gene can also cause the adult-onset form of vitelliform macular dystrophy; however, less than a quarter of all people with this form of the condition have mutations in the BEST1 or PRPH2 gene. In most cases, the cause of the adult-onset form is unknown.

The BEST1 gene provides instructions for making a protein called bestrophin. This protein acts as a channel that controls the movement of charged chlorine atoms (chloride ions) into or out of cells in the retina. Mutations in the BEST1 gene probably lead to the production of an abnormally shaped channel that cannot properly regulate the flow of chloride. Researchers have not determined how these malfunctioning channels are related to the buildup of lipofuscin in the macula and progressive vision loss.
The *PRPH2* gene provides instructions for making a protein called peripherin 2. This protein is essential for the normal function of light-sensing (photoreceptor) cells in the retina. Mutations in the *PRPH2* gene cause vision loss by disrupting structures in these cells that contain light-sensing pigments. It is unclear why *PRPH2* mutations affect only central vision in people with adult-onset vitelliform macular dystrophy.

**Inheritance Pattern**

Best disease is inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to cause the disorder. In most cases, an affected person has one parent with the condition.

The inheritance pattern of adult-onset vitelliform macular dystrophy is uncertain. Some studies have suggested that this disorder may be inherited in an autosomal dominant pattern. It is difficult to be sure, however, because many affected people have no history of the disorder in their family, and only a small number of affected families have been reported.

**Other Names for This Condition**

- vitelliform dystrophy

**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=%22vitelliform+macular+dystrophy%22

**Other Diagnosis and Management Resources**

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Macula (image)
  https://medlineplus.gov/ency/imagepages/9608.htm
- Health Topic: Retinal Disorders
  https://medlineplus.gov/retinaldisorders.html

Genetic and Rare Diseases Information Center

- Adult-onset vitelliform macular dystrophy
- Best vitelliform macular dystrophy
- Macular dystrophy, atypical vitelliform

Additional NIH Resources

- National Eye Institute: Diagram of the Eye
  https://nei.nih.gov/health/eyediagram/

Educational Resources

- MalaCards: best vitelliform macular dystrophy
  https://www.malacards.org/card/best_vitelliform_macular_dystrophy
- Orphanet: Best vitelliform macular dystrophy
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=1243

Patient Support and Advocacy Resources

- Foundation Fighting Blindness
  https://www.fightingblindness.org/diseases/best-disease
- Macular Degeneration Foundation
  https://eyesight.org/
- National Organization for Rare Disorders (NORD)
  https://rarediseases.org/rare-diseases/best-vitelliform-macular-dystrophy/
- Ophthalmic Edge
  https://ophthalmicedge.org/patient/
- Retina International
  http://www.retina-international.org/for-patients/rare-conditions/what-is-best-disease/
Clinical Information from GeneReviews

- Best Vitelliform Macular Dystrophy
  https://www.ncbi.nlm.nih.gov/books/NBK1167

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28vitelliform+macular+dystrophy%5BTIAB%5D%29+OR+%28vitelliform+dystrophy%5BTIAB%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1080+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- MACULAR DYSTROPHY, VITELLIFORM, 2
  http://omim.org/entry/153700
- MACULAR DYSTROPHY, VITELLIFORM, 3
  http://omim.org/entry/608161

Medical Genetics Database from MedGen

- Juvenile Onset Vitelliform Macular Dystrophy
- Vitelliform macular dystrophy
- Vitelliform macular dystrophy type 2

Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19375515
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  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/10854112

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Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/10737974

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