



Multiple cutaneous and mucosal venous malformations

Multiple cutaneous and mucosal venous malformations (also known as VMCM) are bluish patches (lesions) on the skin (cutaneous) and the mucous membranes, such as the lining of the mouth and nose. These lesions represent areas where the underlying veins and other blood vessels did not develop properly (venous malformations). The lesions can be painful, especially when they extend from the skin into the muscles and joints, or when a calcium deposit forms within the lesion causing inflammation and swelling.

Most people with VMCM are born with at least one venous malformation. As affected individuals age, the lesions present from birth usually become larger and new lesions often appear. The size, number, and location of venous malformations vary among affected individuals, even among members of the same family.

Frequency

VMCM appears to be a rare disorder, although its prevalence is unknown.

Causes

Mutations in the *TEK* gene (also called the *TIE2* gene) cause VMCM. The *TEK* gene provides instructions for making a protein called TEK receptor tyrosine kinase. This receptor protein triggers chemical signals needed for forming blood vessels (angiogenesis) and maintaining their structure. This signaling process facilitates communication between two types of cells within the walls of blood vessels, endothelial cells and smooth muscle cells. Communication between these two cell types is necessary to direct angiogenesis and ensure the structure and integrity of blood vessels.

TEK gene mutations that cause VMCM result in a TEK receptor that is always turned on (overactive). An overactive TEK receptor is thought to disrupt the communication between endothelial cells and smooth muscle cells. It is unclear how a lack of communication between these cells causes venous malformations. These abnormal blood vessels show a deficiency of smooth muscle cells while endothelial cells are maintained. Venous malformations cause lesions below the surface of the skin or mucous membranes, which are characteristic of VMCM.

Inheritance Pattern

VMCM is inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to increase the risk of developing venous malformations.

Some gene mutations are acquired during a person's lifetime and are present only in certain cells. These changes, which are not inherited, are called somatic mutations. Researchers have discovered that some VMCM lesions have one inherited and one somatic *TEK* gene mutation. It is not known if the somatic mutation occurs before or after the venous malformation forms. As lesions are localized and not all veins are malformed, it is thought that the inherited mutation alone is not enough to cause venous malformations.

In most cases, an affected person has one parent with the condition.

Other Names for This Condition

- mucocutaneous venous malformations
- VMCM
- VMCM1

Diagnosis & Management

Genetic Testing Information

- What is genetic testing?
[/primer/testing/genetic-testing](#)
- Genetic Testing Registry: Multiple Cutaneous and Mucosal Venous Malformations
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C1838437/>

Other Diagnosis and Management Resources

- GeneReview: Multiple Cutaneous and Mucosal Venous Malformations
<https://www.ncbi.nlm.nih.gov/books/NBK1967>

Additional Information & Resources

Health Information from MedlinePlus

- Health Topic: Skin Conditions
<https://medlineplus.gov/skinconditions.html>
- Health Topic: Vascular Diseases
<https://medlineplus.gov/vasculardiseases.html>

Educational Resources

- Boston Children's Hospital: Venous Malformation in Children
<http://www.childrenshospital.org/conditions-and-treatments/conditions/v/venous-malformation>
- Cincinnati Children's Hospital: Venous Malformations
<https://www.cincinnatichildrens.org/health/v/venous>
- Orphanet: Mucocutaneous venous malformations
https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=2451

Patient Support and Advocacy Resources

- University of Kansas Medical Center Resource List: Dermatology and Genetics
<http://www.kumc.edu/gec/support/derm.html>

Clinical Information from GeneReviews

- Multiple Cutaneous and Mucosal Venous Malformations
<https://www.ncbi.nlm.nih.gov/books/NBK1967>

Scientific Articles on PubMed

- PubMed
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28venous+malformations%5BTIAB%5D%29+AND+%28TIE2%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D>

Catalog of Genes and Diseases from OMIM

- VENOUS MALFORMATIONS, MULTIPLE CUTANEOUS AND MUCOSAL
<http://omim.org/entry/600195>

Sources for This Summary

- Brouillard P, Vikkula M. Genetic causes of vascular malformations. *Hum Mol Genet.* 2007 Oct 15;16 Spec No. 2:R140-9. Epub 2007 Jul 31. Review.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/17670762>
- Limaye N, Boon LM, Vikkula M. From germline towards somatic mutations in the pathophysiology of vascular anomalies. *Hum Mol Genet.* 2009 Apr 15;18(R1):R65-74. doi: 10.1093/hmg/ddp002. Review.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/19297403>
Free article on PubMed Central: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2657941/>
- Limaye N, Wouters V, Uebelhoer M, Tuominen M, Wirkkala R, Mulliken JB, Eklund L, Boon LM, Vikkula M. Somatic mutations in angiopoietin receptor gene TEK cause solitary and multiple sporadic venous malformations. *Nat Genet.* 2009 Jan;41(1):118-24. doi: 10.1038/ng.272. Epub 2008 Dec 14.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/19079259>
Free article on PubMed Central: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2670982/>

- Morris PN, Dunmore BJ, Tadros A, Marchuk DA, Darland DC, D'Amore PA, Brindle NP. Functional analysis of a mutant form of the receptor tyrosine kinase Tie2 causing venous malformations. *J Mol Med (Berl)*. 2005 Jan;83(1):58-63. Epub 2004 Oct 29.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/15526080>
 - Vikkula M, Boon LM, Carraway KL 3rd, Calvert JT, Diamonti AJ, Goumnerov B, Pasyk KA, Marchuk DA, Warman ML, Cantley LC, Mulliken JB, Olsen BR. Vascular dysmorphogenesis caused by an activating mutation in the receptor tyrosine kinase TIE2. *Cell*. 1996 Dec 27;87(7):1181-90.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/8980225>
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Reprinted from Genetics Home Reference:

<https://ghr.nlm.nih.gov/condition/multiple-cutaneous-and-mucosal-venous-malformations>

Reviewed: August 2009

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