Malignant hyperthermia

Malignant hyperthermia is a severe reaction to particular anesthetic drugs that are often used during surgery and other invasive procedures. Specifically, this reaction occurs in response to some anesthetic gases, which are used to block the sensation of pain, either alone or in combination with a muscle relaxant that is used to temporarily paralyze a person during a surgical procedure. If given these drugs, people at risk of malignant hyperthermia may experience a rapid increase in heart rate and body temperature (hyperthermia), abnormally fast breathing, muscle rigidity, breakdown of muscle fibers (rhabdomyolysis), and increased acid levels in the blood and other tissues (acidosis). Without prompt treatment and cessation of the drugs, the body's reaction can cause multiple organs to be unable to function, including the heart (cardiac arrest) and kidneys (renal failure), and it can cause a blood clotting abnormality called disseminated intravascular coagulation. These complications may be life-threatening. (In medicine, the term malignant refers to conditions that are dangerous to one's health.)

People at increased risk of this disorder are said to have malignant hyperthermia susceptibility. Affected individuals may never know they have the condition unless they have a severe reaction to anesthesia during a surgical procedure or they undergo testing (for instance, if susceptibility is suspected because a family member had a severe reaction). Malignant hyperthermia may not occur every time anesthesia is used. Many individuals who develop a severe reaction have previously been exposed to a triggering drug and not had a reaction.

Affected individuals may be at increased risk for "awake" malignant hyperthermia, in which the severe reaction occurs in response to physical activity, often while sick, rather than in reaction to exposure to a triggering drug.

While malignant hyperthermia often occurs in people without other serious medical problems, certain inherited muscle diseases (including central core disease, multiminicore disease, and STAC3 disorder) are associated with malignant hyperthermia susceptibility.

Frequency

Malignant hyperthermia occurs in 1 in 5,000 to 50,000 instances in which people are given anesthetic gases. Susceptibility to malignant hyperthermia is probably more frequent, because many people with an increased risk of this condition are never exposed to drugs that would trigger a reaction and bring them to medical attention.
Causes
Certain variations of the \textit{RYR1} and \textit{CACNA1S} genes increase the risk of developing malignant hyperthermia. Mutations in the \textit{RYR1} gene account for most cases of malignant hyperthermia susceptibility, while mutations in the \textit{CACNA1S} gene cause less than 1 percent of all cases of malignant hyperthermia susceptibility.

The \textit{RYR1} and \textit{CACNA1S} genes provide instructions for making proteins that play essential roles in muscles used for movement (skeletal muscles). For the body to move normally, these muscles must tense (contract) and relax in a coordinated way. Muscle contractions are triggered by the flow of certain charged atoms (ions) in muscle cells. The proteins produced from the \textit{RYR1} and \textit{CACNA1S} genes are involved in the movement of calcium ions within muscle cells. In response to certain signals, the CACNA1S protein helps turn on (activate) the \textit{RYR1} channel. When the \textit{RYR1} channel is active, it releases calcium ions from storage into the fluid-filled space inside muscle cells. The resulting increase in calcium ion concentration inside the cells stimulates muscles to contract.

Mutations in the \textit{RYR1} or \textit{CACNA1S} gene cause the \textit{RYR1} channel to open more easily and close more slowly in response to certain drugs. As a result, abnormally large amounts of calcium ions are released from storage within muscle cells. The abnormal increase in calcium ion concentration within muscle cells activates processes that generate heat (leading to increased body temperature) and produce excess acid (leading to acidosis). An overabundance of calcium ions also causes skeletal muscles to contract, which leads to muscle rigidity.

Up to half of people with malignant hyperthermia susceptibility do not have a mutation in one of the known genes. The causes of these cases are still under study.

Inheritance Pattern
Malignant hyperthermia susceptibility 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 a severe reaction to certain drugs used during surgery. In most cases, an affected person inherits the altered gene from a parent who is also at risk for the condition.

Other Names for This Condition
- anesthesia related hyperthermia
- hyperpyrexia, malignant
- hyperthermia, malignant
- malignant hyperthermia
- MHS
Diagnosis & Management

Genetic Testing Information

- What is genetic testing?
  /primer/testing/genetictesting

- Genetic Testing Registry: Malignant hyperthermia susceptibility type 2

- Genetic Testing Registry: Malignant hyperthermia susceptibility type 3

- Genetic Testing Registry: Malignant hyperthermia susceptibility type 4

- Genetic Testing Registry: Malignant hyperthermia susceptibility type 5

- Genetic Testing Registry: Malignant hyperthermia susceptibility type 6

- Genetic Testing Registry: Malignant hyperthermia, susceptibility to, 1

Research Studies from ClinicalTrials.gov

- ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22malignant+hyperthermia%22

Other Diagnosis and Management Resources

- GeneReview: Malignant Hyperthermia Susceptibility
  https://www.ncbi.nlm.nih.gov/books/NBK1146

- MedlinePlus Encyclopedia: Malignant Hyperthermia
  https://medlineplus.gov/ency/article/001315.htm

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Malignant Hyperthermia
  https://medlineplus.gov/ency/article/001315.htm

- Health Topic: Muscle Disorders
  https://medlineplus.gov/muscledisorders.html
Genetic and Rare Diseases Information Center

- King Denborough syndrome
- Malignant hyperthermia
  https://rarediseases.info.nih.gov/diseases/6964/malignant-hyperthermia

Educational Resources

- JAMA Patient Page: Malignant Hyperthermia
  https://jamanetwork.com/journals/jama/fullarticle/201084
- MalaCards: malignant hyperthermia
  https://www.malacards.org/card/malignant_hyperthermia
- Orphanet: Malignant hyperthermia of anesthesia
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=423

Patient Support and Advocacy Resources

- Malignant Hyperthermia Association of the United States
  https://www.mhaus.org/
- National Organization for Rare Disorders (NORD): Malignant Hyperthermia
  https://rarediseases.org/rare-diseases/malignant-hyperthermia/
- National Organization for Rare Disorders (NORD): RYR-1-Related Diseases
  https://rarediseases.org/rare-diseases/ryr-1-related-diseases/
- North American Malignant Hyperthermia Registry
  https://anest.ufl.edu/namhr/
- Resource List from the University of Kansas Medical Center
  http://www.kumc.edu/gec/support/malighyp.html

Clinical Information from GeneReviews

- Malignant Hyperthermia Susceptibility
  https://www.ncbi.nlm.nih.gov/books/NBK1146

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28Malignant+Hyperthermia%5BMAJR%5D%29+AND+%28malignant+hyperthermia%5BTIAB%5D%29+AND+en glish%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+720+days%22%5Bdp %5D
Catalog of Genes and Diseases from OMIM

- MALIGNANT HYPERTERMIA, SUSCEPTIBILITY TO, 1
  http://omim.org/entry/145600

- MALIGNANT HYPERTERMIA, SUSCEPTIBILITY TO, 2
  http://omim.org/entry/154275

- MALIGNANT HYPERTERMIA, SUSCEPTIBILITY TO, 3
  http://omim.org/entry/154276

- MALIGNANT HYPERTERMIA, SUSCEPTIBILITY TO, 4
  http://omim.org/entry/600467

- MALIGNANT HYPERTERMIA, SUSCEPTIBILITY TO, 5
  http://omim.org/entry/601887

- MALIGNANT HYPERTERMIA, SUSCEPTIBILITY TO, 6
  http://omim.org/entry/601888

Sources for This Summary


   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/30168660

   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/30805902

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