

# **Anencephaly**

# **Description**

Anencephaly is a condition that prevents the normal development of the brain and the bones of the skull. This condition results when a structure called the neural tube fails to close during the first few weeks of embryonic development. The neural tube is a layer of cells that ultimately develops into the brain and spinal cord. Because anencephaly is caused by abnormalities of the neural tube, it is classified as a neural tube defect.

Because the neural tube fails to close properly, the developing brain and spinal cord are exposed to the amniotic fluid that surrounds the fetus in the womb. This exposure causes the nervous system tissue to break down (degenerate). As a result, people with anencephaly are missing large parts of the brain called the cerebrum and cerebellum. These brain regions are necessary for thinking, hearing, vision, emotion, and coordinating movement. The bones of the skull are also missing or incompletely formed.

Because these nervous system abnormalities are so severe, almost all babies with an encephaly die before birth or within a few hours or days after birth.

# Frequency

Anencephaly is one of the most common types of neural tube defect, affecting about 1 in 1,000 pregnancies. However, most of these pregnancies end in miscarriage, so the prevalence of this condition in newborns is much lower. An estimated 1 in 10,000 infants in the United States is born with anencephaly.

#### Causes

Anencephaly is a complex condition that is likely caused by the interaction of multiple genetic and environmental factors. Some of these factors have been identified, but many remain unknown.

Changes in dozens of genes in individuals with anencephaly and in their mothers may influence the risk of developing this type of neural tube defect. The best-studied of these genes is *MTHFR*, which provides instructions for making a protein that is involved in processing the vitamin folate (also called vitamin B9). While a shortage (deficiency) of this vitamin is an established risk factor for neural tube defects, there are many factors that can contribute to folate deficiency. Changes in other genes related to folate processing and genes involved in the development of the neural tube have also been

studied as potential risk factors for an encephaly. However, no genes appear to play a major role in causing the condition.

Researchers have also examined environmental factors that could contribute to the risk of anencephaly. Folate deficiency plays a significant role. Studies have shown that women who take supplements containing folic acid (the synthetic form of folate) before they get pregnant and very early in their pregnancy are significantly less likely to have a baby with a neural tube defect, including anencephaly. Other possible maternal risk factors for anencephaly include diabetes mellitus, obesity, exposure to high heat (such as a fever or use of a hot tub or sauna) in early pregnancy, and the use of certain anti-seizure medications during pregnancy. However, it is unclear how these factors may influence the risk of anencephaly.

# Learn more about the gene associated with Anencephaly

MTHFR

### Inheritance

Most cases of anencephaly are sporadic, which means they occur in people with no history of the disorder in their family. A small percentage of cases have been reported to run in families; however, the condition does not have a clear pattern of inheritance. For parents who have had a child with anencephaly, the risk of having another affected child is increased compared with the risk in the general population.

## Other Names for This Condition

- Anencephalia
- Anencephalus
- Aprosencephaly
- Congenital absence of brain

#### Additional Information & Resources

## **Genetic Testing Information**

- Genetic Testing Registry: Anencephaly (https://www.ncbi.nlm.nih.gov/gtr/conditions/ C0002902/)
- Genetic Testing Registry: Neural tube defect (https://www.ncbi.nlm.nih.gov/gtr/conditions/C0027794/)
- Genetic Testing Registry: Neural tube defects, folate-sensitive (https://www.ncbi.nlm.nih.gov/qtr/conditions/C1866558/)

## Genetic and Rare Diseases Information Center

Isolated anencephaly/exencephaly (https://rarediseases.info.nih.gov/diseases/5808/index)

## Patient Support and Advocacy Resources

National Organization for Rare Disorders (NORD) (https://rarediseases.org/)

## Clinical Trials

ClinicalTrials.gov (https://clinicaltrials.gov/search?cond=%22Anencephaly%22)

## Catalog of Genes and Diseases from OMIM

- NEURAL TUBE DEFECTS, FOLATE-SENSITIVE; NTDFS (https://omim.org/entry/6 01634)
- NEURAL TUBE DEFECTS, SUSCEPTIBILITY TO; NTD (https://omim.org/entry/182 940)
- ANENCEPHALY 1; ANPH1 (https://omim.org/entry/206500)

#### Scientific Articles on PubMed

 PubMed (https://pubmed.ncbi.nlm.nih.gov/?term=%28Anencephaly%5BMAJR%5D %29+AND+%28anencephaly%5BTIAB%5D%29+AND+english%5Bla%5D+AND+hu man%5Bmh%5D+AND+%22last+1800+days%22%5Bdp%5D)

#### References

- Au KS, Ashley-Koch A, Northrup H. Epidemiologic and genetic aspects of spinabifida and other neural tube defects. Dev Disabil Res Rev. 2010;16(1):6-15. doi: 10.1002/ddrr.93. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/20419766) or Free article on PubMed Central (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC305 3142/)
- Bassuk AG, Kibar Z. Genetic basis of neural tube defects. Semin PediatrNeurol. 2009 Sep;16(3):101-10. doi: 10.1016/j.spen.2009.06.001. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/19778707)
- Botto LD, Moore CA, Khoury MJ, Erickson JD. Neural-tube defects. N Engl J Med. 1999 Nov 11;341(20):1509-19. doi: 10.1056/NEJM199911113412006. No abstractavailable. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/10559453)
- Copp AJ, Greene ND. Genetics and development of neural tube defects. J Pathol. 2010 Jan;220(2):217-30. doi: 10.1002/path.2643. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/19918803) or Free article on PubMed Central (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4239538/)

- Doudney K, Grinham J, Whittaker J, Lynch SA, Thompson D, Moore GE, Copp AJ, Greene ND, Stanier P. Evaluation of folate metabolism gene polymorphisms as riskfactors for open and closed neural tube defects. Am J Med Genet A. 2009Jul; 149A(7):1585-9. doi: 10.1002/ajmg.a.32937. No abstract available. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/19533788)
- Greene ND, Stanier P, Copp AJ. Genetics of human neural tube defects. Hum MolGenet. 2009 Oct 15;18(R2):R113-29. doi: 10.1093/hmg/ddp347. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/19808787) or Free article on PubMed Central (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2758708/)
- Hickey SE, Curry CJ, Toriello HV. ACMG Practice Guideline: lack of evidencefor MTHFR polymorphism testing. Genet Med. 2013 Feb;15(2):153-6. doi:10.1038/gim. 2012.165. Epub 2013 Jan 3. Erratum In: Genet Med. 2020Dec;22(12):2125. doi: 10. 1038/s41436-020-0843-0. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/232 88205)
- Levin BL, Varga E. MTHFR: Addressing Genetic Counseling Dilemmas
   UsingEvidence-Based Literature. J Genet Couns. 2016 Oct;25(5):901-11. doi:10.
   1007/s10897-016-9956-7. Epub 2016 Apr 30. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/27130656)
- Obeidi N, Russell N, Higgins JR, O'Donoghue K. The natural history ofanencephaly. Prenat Diagn. 2010 Apr;30(4):357-60. doi: 10.1002/pd.2490. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/20198650)
- Wilson RD; SOGC GENETICS COMMITTEE; SPECIAL CONTRIBUTOR. RETIRED: Prenatalscreening, diagnosis, and pregnancy management of fetal neural tube defects. JObstet Gynaecol Can. 2014 Oct;36(10):927-939. doi: 10.1016/S1701-2163(15)30444-8. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/25375307)
- Yan L, Zhao L, Long Y, Zou P, Ji G, Gu A, Zhao P. Association of the maternalMTHFR C677T polymorphism with susceptibility to neural tube defects inoffsprings: evidence from 25 case-control studies. PLoS One. 2012;7(10):e41689. doi: 10.1371/journal.pone.0041689. Epub 2012 Oct 3. Citation on PubMed (https://p ubmed.ncbi.nlm.nih.gov/23056169) or Free article on PubMed Central (https://www. ncbi.nlm.nih.gov/pmc/articles/PMC3463537/)
- Zhang T, Lou J, Zhong R, Wu J, Zou L, Sun Y, Lu X, Liu L, Miao X, Xiong G.
  Genetic variants in the folate pathway and the risk of neural tube defects: ametaanalysis of the published literature. PLoS One. 2013 Apr 4;8(4):e59570. doi:10.1371/
  journal.pone.0059570. Print 2013. Citation on PubMed (https://pubmed.ncbi.nlm.nih.
  gov/23593147) or Free article on PubMed Central (https://www.ncbi.nlm.nih.gov/pmc/
  articles/PMC3617174/)

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