Anencephaly [1]


Anencephaly is an open neural tube [5] defect, meaning that part of the neural tube [5] does not properly close or that it has reopened during early embryogenesis [6]. An embryo with anencephaly develops without the top of the skull, but retains a partial skull, including the face. Anencephaly is one of the most common birth defects [7] of the neural tube [5], occurring at a rate of approximately one in one thousand human pregnancies. The condition can be caused by environmental exposure to chemicals, dietary deficiencies, or genetic mutations.


In addition to nutritional causes, certain genetic abnormalities potentially cause anencephaly. Mutations in genes [15] such as MacMARCKS, which is active during neural tube [5] development, and FOXN1, a gene that is important for the proper development of the central nervous system [16], have been suggested to cause anencephaly. A 1996 study suggested that the protein coded by the MacMARCKS gene may play a role in closing the neural tube [5]. The MacMARCKS protein embeds in the cellular membrane that regulates binding of actin filaments between cells. Jianmin Chen and colleagues at The Rockefeller University [17] in New York, New York, examined mouse [18] embryos with normal MacMARCKS gene expression patterns and noted that the brain developed normally with no cases of anencephaly or other identifiable neural tube [5] defects. They then created mutants which expressed a nonfunctional MacMARCKS protein, and they found that all mutant embryos were anencephalic. Upon further analysis, researchers found the wild type MacMARCKS protein in the area of neural tube [5] closure, concluding that MacMARCKS plays an important role in the proper closure of the neural tube [5].

In 2007 Stefania Amorosi and colleagues, at the University of Naples in Naples, Italy, examined anencephalic mouse [18] embryos and an anencephalic human embryo. The embryos were homozygous for a mutation in the FOXN1 gene. Amorosi found that the FOXN1 gene was expressed in the choroid plexus of mice, a structure in the ventricles of the mammalian brain responsible for secreting cerebrospinal fluid (CSF). CSF is a tissue that transports signaling molecules and directs brain development. The FOXN1 gene codes for a transcription factor protein that controls the expression of genes [15] that contribute to development of the central nervous system [16]. Amorosi and colleagues concluded that mutations in FOXN1 resulted in improper neural tube [5] formation, leading to anencephaly.

There is also evidence that environmental exposure to certain chemicals early in pregnancy [9] can cause anencephaly. A 2001 study by Julia Blanco Muñoz, at the University of Granada in Granada, Spain, surveyed the families of 157 Mexican children born with anencephaly and 151 families whose children developed normally. The study showed that mothers exposed to pesticides such as methyl parathion during their first month of pregnancy [9] were five times more likely to give birth to a child with anencephaly than were women not exposed to pesticides. The study also showed that families were twice as likely to have an anencephalic child when the father was exposed to methyl parathion at any time prior to conception [19]. As of 2002 Mexico had the world’s highest rate of anencephaly, with 8.05 cases per 10,000 live births.
Anencephaly can be diagnosed using a sonogram, an imaging method using ultrasound technology to visualize the fetus inside the mother. An anencephalic fetus has a triangular-shaped face and lacks the anterior region of the skull, both of which can be seen on the sonogram. Using a sonogram, anencephaly can be diagnosed as early as eleven weeks into the pregnancy. An older method for diagnosing anencephaly is using karyotyping, a method of staining chromosomes with dye, and investigating chromosome structure with a microscope. Researchers have found evidence of a connection between anencephaly and chromosome 13, when the latter has the shape of a ring. In these cases, genes which are responsible for proper brain development are deleted, and the remaining sections of the chromosome bind at the ends, forming a ring.

Some controversy accompanies the idea of using anencephalic fetuses as organ donors. Anencephalic fetuses are unable to survive outside of the womb, but they develop many functional organs, and anencephalic fetuses are candidates for organ donation. Proponents of embryonic organ donation state that because large sections of the brain are missing, anencephalic fetuses cannot feel pain and are never conscious. Ethicists, scientists, and doctors grapple with questions about the morality of distributing organs from anencephalic fetuses.

Sources


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