



The Role of Genetic Mutations on Genes PAX2 & PAX6 in Cyclopia Syndrome

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Review Article

Volume 5 Issue 1

Received Date: March 02, 2020

Published Date: May 28, 2020

DOI: 10.23880/nnoj-16000147

Abstract

Cyclopia syndrome typically coexists with holoprosencephaly. This ocular lesion may present as complete ocular fusion in a single orbit or as two eyes in a single orbit. Eye defects linked with cyclopia syndrome include colobomas (gaps) in the iris, retina, and optic nerve; inconstant optic nerve numbers (either one or two is possible), and an absent or abnormal optic chiasm. Cyclopia syndrome is caused by mutations in the PAX2 and PAX6 genes. Cyclopia syndrome occurs when the rostral (anterior) portion of the notochord and adjacent mesoderm are deficient in mass. This shortage leads to the aberrant induction of the forebrain tissues followed by severe derangement of midline facial development.

Keywords: Cyclopia Syndrome; Genetic Mutations; PAX2; PAX6 Genes; Holoprosencephaly

Overview of Cyclopia Syndrome

Cyclopia syndrome is a rare disorder of holoprosencephaly (fracture of the brain in the fetus) that occurs with birth defects. In this disease, the orbit of the eye is divided into two very close caverns, with either the monocular organism being seen or the binoculars being closely related.

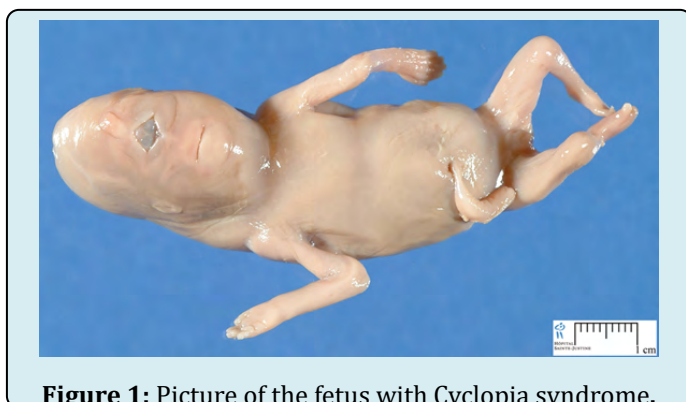


Figure 1: Picture of the fetus with Cyclopia syndrome.

The syndrome affects both animals and humans, with the frequency in animals 1 in every 16,000 births and in the human fetus 1 in 250 births [1].

Signs and Symptoms of Cyclopia Syndrome

Typically, patients with cyclopia syndrome have a long nose or inadequate nose in the form of a long snout. Patients with cyclopia syndrome have one eye in the middle of the forehead or two eyes completely clinging. The ears are bent or fractured or crumpled inward, and most patients have a fleshy appendage 3 to 5 cm in the middle of their forehead [1].

Cyclopia syndrome most commonly involves embryos that are either aborted or born at the same age as the fetus and, in good fortune, will be certain shortly after their death. Cyclopia syndrome is a rare genetic disorder that has been reported in only a few cases in the medical world, and most are preserved at the vrolijk Amsterdam Museum of Medicine

in the Netherlands [1].

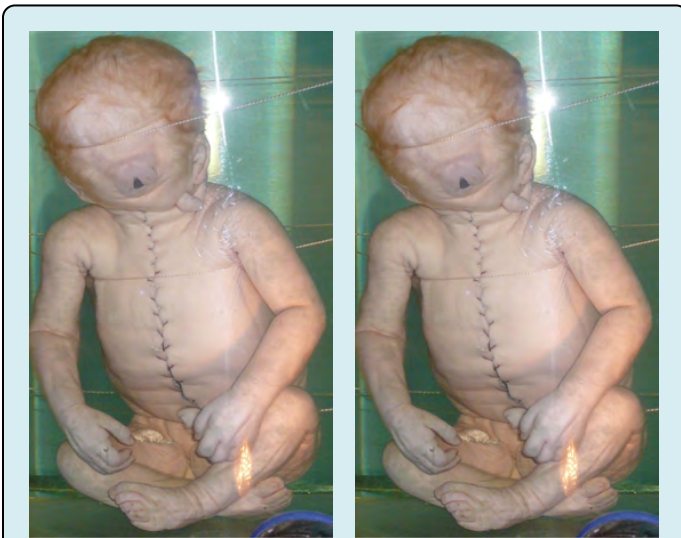


Figure 2: Image of human with cyclopia syndrome with indicative features.

Animals that develop cyclopia syndrome include farm animals such as horses, sheep, pigs, and chickens. In these animals, like humans, the nasal organs form inefficiently, which shortly after birth causes the airway to pass through the respiratory tract, resulting in suffocation [1,2].

Etiology of Cyclopia Syndrome

One of the major causes of cyclopia syndrome can be increased maternal age during pregnancy and the susceptibility of cytopenia to cyclopia syndrome in post-fertilization and zygote cell formation, as well as the use of toxic or sedative alkaloids. Maternal pregnancy can increase the susceptibility of the fetus to cyclopia syndrome. In fact, most people with cyclopia syndrome are due to the mother's mistake during pregnancy in using herbs such as veratrum californicum to improve vomiting and cramping and poor

blood circulation. In other words, this plant has alkaloids, although it is known as a natural painkiller in general. Of course, the use of this medicinal plant during pregnancy has no adverse effect on women and men, but it was very dangerous during pregnancy due to its alkaloid properties and is a potent teratogenic agent. This alkaloid property of the plant disturbs the mitotic and spindle filament cycles, resulting in the failure of the eye, nose, and ear cavity to form successfully, as well as the correct shape of the brain [1,3].



Figure 3: Picture of Veratrum medicinal plant.

Molecular Genetics of Cyclopia Syndrome

SHH gene, as the regulator of expression and synthesis of proteins of the nervous system for the formation of the brain and PAX6 gene, is the correct state of synthesis of the eyes which, due to chromosomal abnormalities, are mutated and lose their normal function. The SHH gene is located on the long arm of chromosome 7 at 7q36.3, and in addition to the above, the gene regulates the correct shape of the hands and nose. The SHH gene plays a key role in initiating the expression of other genes such as PAX6 and PAX2 in the cell [1,4].

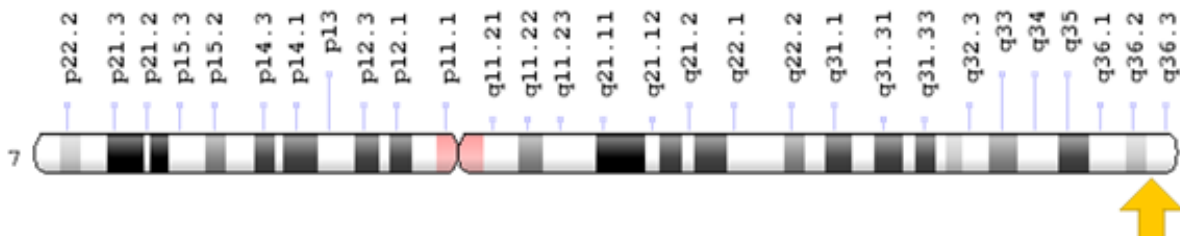
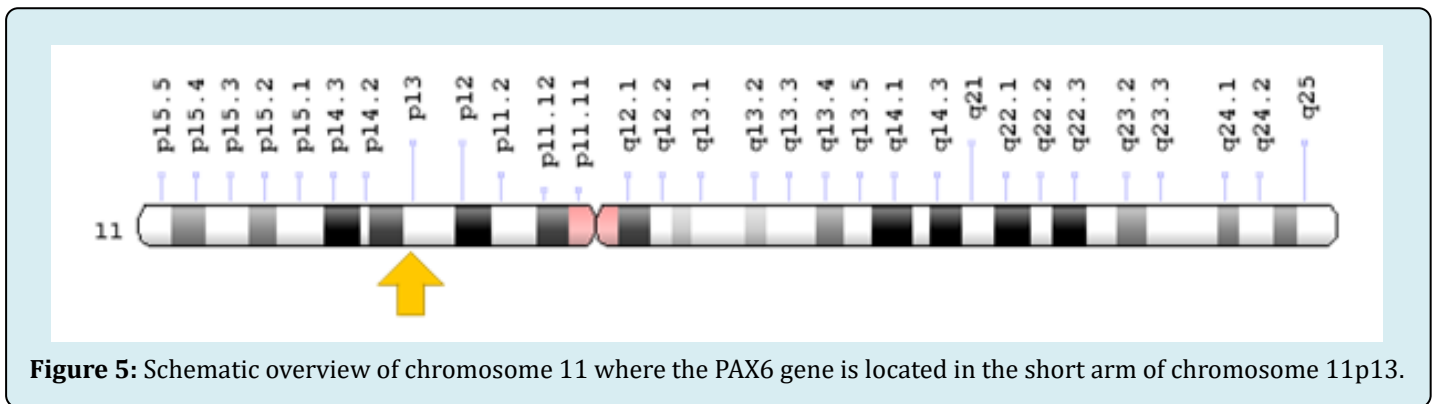
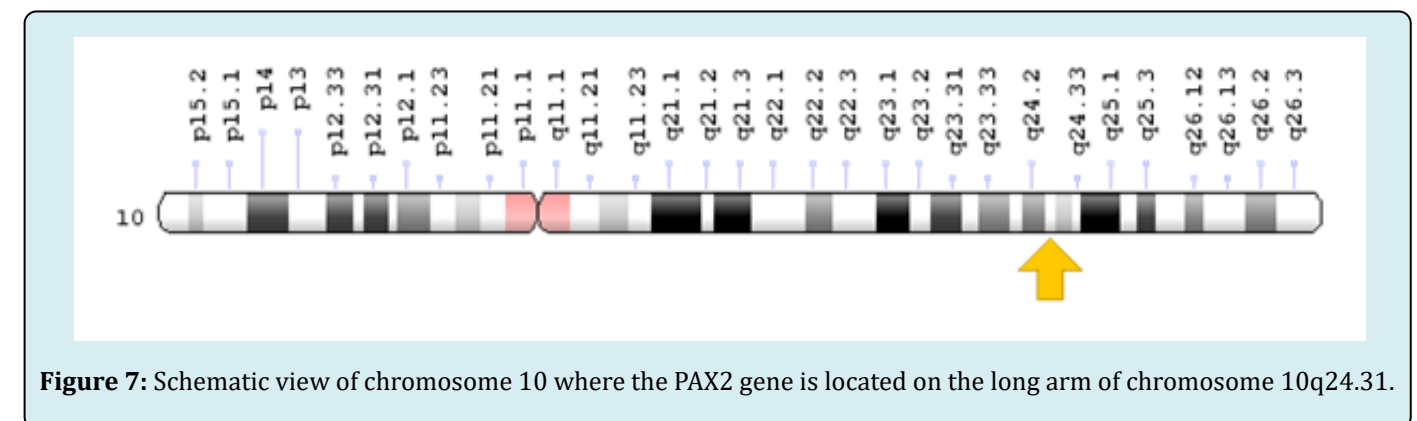
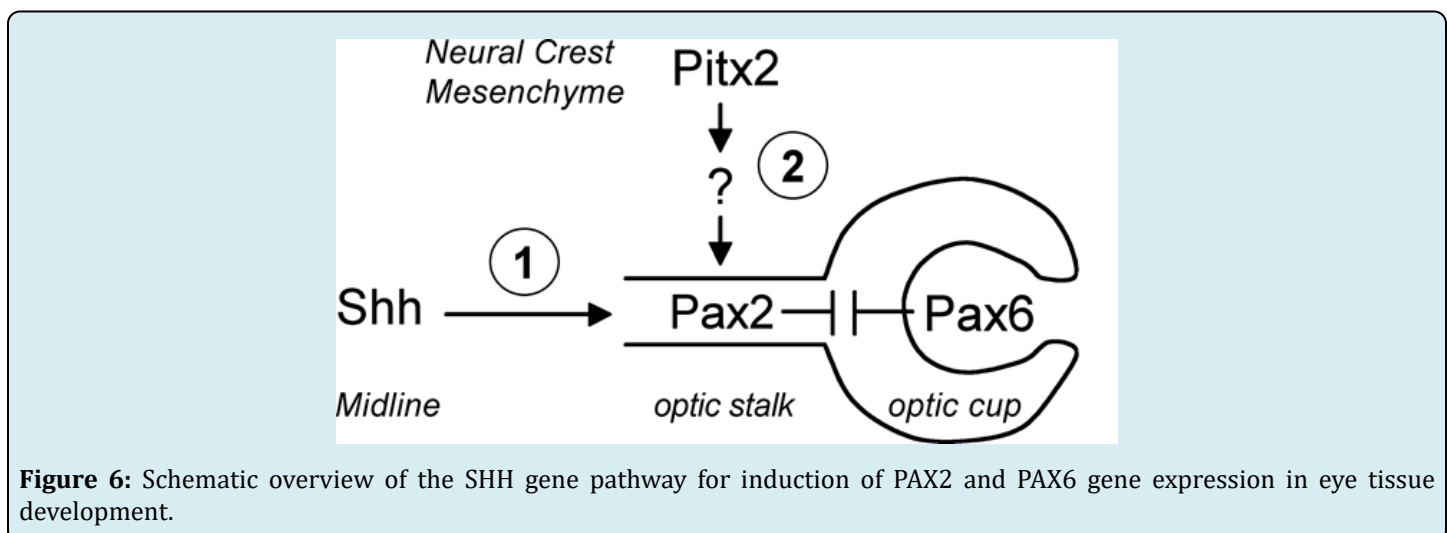


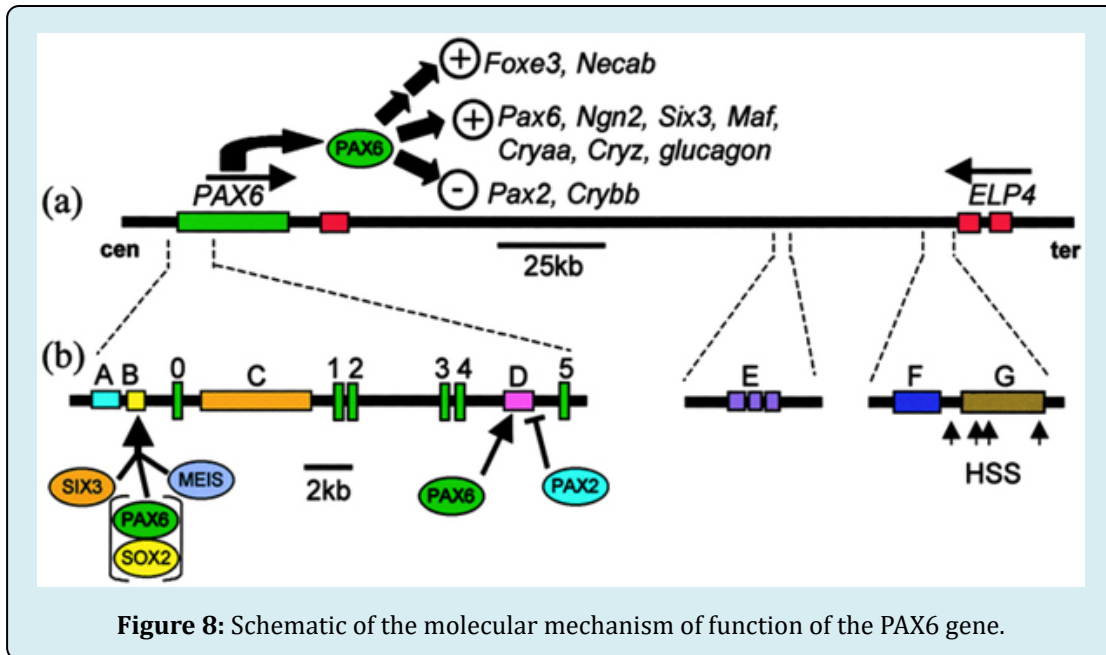
Figure 4: Schematic overview of chromosome 7 where the SHH gene is located in the long arm of chromosome 7q36.3.



The PAX6 gene acts as a natural protein for the eyes. The gene is located on the short arm of chromosome 11 in 11p13.

This gene plays a key role in the evolution of the eyes to the natural form, both in humans and animals [1,5].





Any abnormality in this gene, including nucleotide deletion or deletion or insertion, will lead to impaired vision or blindness. Patients with cyclopia syndrome, on the short arm of chromosome 11 in the 11p11 and 11p13 regions, have mutations that occur and cause visual and visual developmental disorders [1,6].

Diagnosis of Cyclopia Syndrome

Cyclopia syndrome is diagnosed based on facial features findings with imaging techniques such as ultrasound. Prenatal diagnosis is also possible by molecular genetic testing of amniocentesis fluid and fetal placenta chorionic villi [1,7].

History of Cyclopia Syndrome

The word cyclops in ancient Greek means an eye circle referring to an athlete in ancient Greek mythology. This hero had one eye in the center of the forehead. The name of this hero was Polyphemus. He was the son of Uranus and Gaia of the great Titans of ancient Greece [1,8].

But the first report of cyclopia syndrome goes back to 1665 on a horse in the UK. Then in March 1793, a 46 - year - old woman in Sweden gave birth to a child with cyclopia syndrome who died after two hours. On December 28, 2005, a kitten in Redmond, USA, was born with cyclopia syndrome, which died a day after giving birth. In 2006, a baby girl in India was born with cyclopia syndrome that died a day after giving birth. In 2011, an embryo with cyclopia syndrome was discovered from the belly of a giant hunting shark in Mexico.

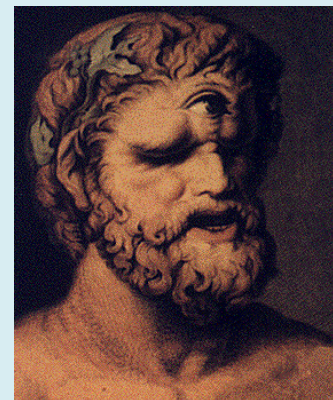


Figure 9: Painting image of the Athenian god Polyphemus, who had an eye in the middle of the forehead.



Figure 10: Image of a baby with cyclopia syndrome.

In 2012, a kitten was born with cyclopia syndrome who died after two hours. Cyclopia syndrome has symptoms similar to patau *syndrome*. But the phenotype of cyclopia Syndrome is much more severe than that of patau *syndrome* and the half-life of cyclopia syndrome patients is shorter than that of patau *syndrome*. The major genes involved in cyclopia syndrome are: SHH, ZIC2, SLX3, and TGIF1 [1,9].

Discussion and Conclusion

Cyclopia (also cyclocephaly or synophthalmia) is a rare form of holoprosencephaly and is a congenital disorder (birth defect) characterised by the failure of the embryonic prosencephalon to properly divide the orbits of the eye into two cavities. It is the severest facial expression of the holoprosencephaly syndrome. Its incidence is 1 in 100 000 in newborns. Typically, the nose is either missing or replaced with a non-functioning nose in the form of a proboscis. Such a proboscis generally appears above the central eye, or on the back, and is characteristic of a form of cyclopia called rhinencephaly or rhinocephaly. Most such embryos are either naturally aborted or are stillborn on delivery. Some cases of cyclopia have been associated with a rare chromosomal condition called 'Patau Syndrome' (trisomy 13). SHH (Sonic Hedgehog Gene Regulator) is involved in the separation of the single eye field into two bilateral fields. Although not proven, it is thought that SHH emitted from the prechordal plate suppresses Pax6 which causes the eye field to divide into two. If the SHH gene is mutated, the result is cyclopia, a single eye in the centre of the face [1-10].

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