

Features of the Clinical Course of Congenital Myopia

Boboev Saidavzal Abdrahmanovich

Samarkand State Medical University

saidavzalbabaev@gmail.com

Received: 2025, 15, Jan

Accepted: 2025, 21, Feb

Published: 2025, 11, Mar

Copyright © 2025 by author(s) and BioScience Academic Publishing. This work is licensed under the Creative Commons Attribution International License (CC BY 4.0).



Open Access

<http://creativecommons.org/licenses/by/4.0/>

Annotation: Congenital myopia is a decrease in visual acuity caused by intrauterine disorders in the structure and function of various parts of the eyeball. The disease develops under the influence of genetic mutations, teratogenic effects and perinatal complications. Visual impairment is detected in children from an early age, often accompanied by astigmatism and amblyopia. Visometry, ultrasound and OCT of the eye, computer accommodation and laser retinotomography are used for diagnostics. Treatment consists of early optical correction of refractive errors, the use of auxiliary drugs and physiotherapy methods.

Keywords: Causes, Pathogenesis, Classification, Symptoms of congenital myopia, Diagnostics, Treatment of congenital myopia, Optical correction, Pharmacotherapy and physiotherapy, Surgical treatment, Prognosis and prevention.

Introduction: Myopia is the most common visual impairment, occurring in 6-8% of primary school children, 25-30% of school graduates and 35-42% of adults. The incidence of congenital visual impairment in infants under 1 year of age is 1.4-4.5%, more than half of these cases have a high degree of myopia - minus 6 diopters. Congenital myopia is often combined with other eye diseases and is difficult to optically correct, therefore it represents a serious problem in pediatric ophthalmology.

Congenital myopia is caused by pathologies of the visual apparatus that are formed in the antenatal and intranatal periods. The disease is multifactorial and is accompanied by various anatomical, functional and refractive disorders. Possible causes of the disease include:

Genetic factors. There are 23 genes responsible for the predisposition to vision disorders located at different loci on chromosome 15. They are not well understood and are probably part of polygenic systems that determine visual acuity. In addition, 54-65% of children with congenital

myopia have one or both parents who are nearsighted.

Pregnancy pathologies. The risk of developing congenital myopia in the fetus increases with early and late toxicosis, the threat of spontaneous abortion, gynecological and extragenital diseases of the mother. Visual impairment is a common consequence of intrauterine TORCH complex infections.

Premature birth. High levels of congenital myopia are common in children with retinopathy of prematurity. The younger the gestational age of the fetus at the time of premature birth, the higher the risk of ophthalmological diseases.

Complicated labor. Congenital myopia occurs in newborns who have suffered birth trauma to the skull or cervical spine, umbilical cord, or prolonged labor. The disease can be a consequence of perinatal hypoxia and encephalopathy.

Research methods and materials: The development of congenital myopia is based on dystrophic processes and stretching of the sclera apparatus in the posterior part of the eye. The eyeball acquires a pathologically elongated shape, which leads to the convergence of rays passing through the lens in front of the retina, which forms a blurred image. Changes in the structure of the eye are accompanied by disturbances in tissue trophism, which exacerbates existing pathologies and leads to retinal atrophy.

Metabolic disorders play an important role in the development of myopia. In patients, the antioxidant protection of red blood cell membranes is reduced and the metabolism of calcium and phosphorus is disturbed. Congenital myopia is also characterized by impaired synthesis of collagen fibers and proteoglycans, so the disease is often found in children with connective tissue pathologies.

Transient myopia is characterized by the normalization of visual acuity at the beginning of the second year of a child's life as a result of the maturation of the ciliary muscle and a decrease in the refractive power of the anterior segment of the eye.

In true myopia, there is a stable or progressive deterioration of vision, which is accompanied by elongation of the eyeball and changes in the fundus.

Results: Young children do not complain of poor vision, so parents should pay attention to indirect signs. A child with myopia constantly squints, blinks rapidly, and wrinkles appear on his forehead when trying to see distant objects. He holds toys close to his eyes and approaches the TV to study the picture more closely. Older children may complain of blurred vision and headaches after visual stress.

Previously, congenital myopia was considered a stable disease - visual acuity in children remained unchanged for many years. However, in recent decades, doctors have increasingly encountered rapidly developing forms of the disease. In this case, the child's vision is constantly deteriorating, so that, without proper correction, already at primary school age, he sees only objects located at a distance of 1-1.5 meters around him.

Congenital high myopia is a disease that, if left untreated, can lead to disability and even blindness in the patient. Possible complications include clouding of the lens (cataracts), destruction of the vitreous body and detachment of its membrane. Sometimes retinal detachment and retinal tears are observed, which occur against the background of prolonged dystrophy in the dentate gyrus.

The severe course of congenital myopia is caused not only by the high rate of refractive errors. In many patients, myopia is accompanied by amblyopia - the syndrome of "lazy eye". Pathology leads to a further deterioration of visual acuity even with relatively stable sizes and proportions of the structures of the eyeball. With amblyopia, the visual function of one eye is suppressed and binocular perception of the surrounding world is impaired.

Another typical complication of congenital myopia is astigmatism. With myopic astigmatism, the

shape of the cornea and lens is distorted, as a result of which light rays are refracted incorrectly, which forms a blurry, distorted image. If timely correction is not carried out, poor vision causes problems in understanding the world, learning and social adaptation, and causes serious limitations in everyday life.

Patients with complaints of poor vision are consulted by a pediatric or adult ophthalmologist. If a clinical examination reveals a decrease in visual acuity, the presence of myopia is not suspected and does not require differential diagnosis. In the second stage, an expanded instrumental examination is indicated to determine the condition of various eye structures and identify concomitant pathologies. The following methods are used:

Visometry. In patients under 3 years of age, the examination is carried out using the technique of preference testing and optokinetic nystagmus recording, depending on the presence of object vision. For children 3 years of age and older and adults, special tables with pictures, rings or letters are used. Visometry is supplemented by an assessment of visual fields - perimetry.

Ultrasound of the eye. Ultrasound examination reveals enlargement of the anterior-posterior segment of the eyeball, measures the transverse diameter of the eye and the acoustic density of the sclera. Ultrasound data exclude accommodation spasm and confirm the presence of anatomical changes that cause myopia.

Accommodation. In patients with progressive myopia, a special diagnostic method is prescribed to determine the decreasing accommodation and objective accommodative response (OAR). An increase in OAR during treatment is considered a favorable prognostic factor.

Optical coherence tomography. Using OCT, specific features of the structure of the retina in congenital myopia are identified: thickening of the neuroepithelium in the center of the macula, a decrease in the thickness of the parafoveal region, changes in the relief of the retina and its pronounced thinning in the paracentral zones.

Discussion: Heidelberg laser retinotomography. The study shows structural anomalies of the optic nerve head: a bent position, oblique entry of the vessels, atrophy of the adjacent chorioretinal tissues. In congenital myopia, the optic disc has a smaller excavation than in acquired myopia. Posterior staphylomas (protrusions of the sclera) are often detected.

Electroretinography. In patients with congenital myopia, the study of the bioelectric activity of the neuroepithelium reveals stationary night blindness, pathologies of the rod and cone systems, and dysfunctions of the outer layers of the retina.

The main goal of treatment is to correct visual acuity, which begins as early as possible after myopia is detected. For this purpose, glasses, contact lenses, and a combination of the two methods are used. For children under 8 years of age, only glasses are prescribed, since small patients cannot correctly put on and remove lenses even with the help of their parents. In the case of concomitant astigmatism, special cylindrical lenses are used.

Conclusion: In old age, soft contact lenses (SCL) and rigid orthokeratological lenses for night correction are prescribed. They provide clear vision in all areas of the visual field, eliminating the need for the child to wear glasses. MKL is an indispensable type of correction when there is a significant difference in refraction between the left and right eyes. Clinical observations confirm that the transition to contact correction allows you to increase corrected visual acuity by 0.1-0.3 points.

Drug therapy for congenital myopia shows limited effectiveness and is therefore prescribed in combination with other methods. Vitamin and mineral complexes, vasodilators and neurometabolic drugs are used. Drugs are taken orally, instilled into the eyes or administered parabolbarly. Physiotherapy is also used: electrophoresis with a solution of calcium chloride, magnetic therapy, acupuncture and electropuncture.

The operation is recommended for stretching the posterior pole of the eye, myopic staphylomas,

damage to the vascular and nervous structures of this area. Patients are prescribed scleroplasty and eye filling using donor sclera to stop retinal dystrophy and prevent irreversible loss of vision. In 90% of patients, myopia can stabilize, in the remaining 10% of cases, the rate of its development slows down by 2 times;

Modern methods of optical correction allow a patient with congenital myopia to have relatively normal vision and live a full life with minimal restrictions. The long-term prognosis is determined by the timely start of treatment, the degree of myopia and the dynamics of its development. Primary prevention of the disease has not been developed. Secondary prevention includes regular examinations of the child by an ophthalmologist to monitor visual acuity.

List of used literature:

1. Myopia. Clinical recommendations of the All-Russian public organization "Union of Ophthalmologists". – 2017.
2. Clinical, functional and biomechanical aspects of the pathogenesis, diagnosis and treatment of congenital myopia: literature review and analysis of own data / GA Markosyan et al.// Russian Pediatric Ophthalmology. – 2016. – No. 3.
3. Ophthalmology. National leadership / SE Avetisov, EA Egorov, LK Moshetova. – 2014.
4. Morphometric and functional characteristics of the macular region in patients with high congenital myopia / EP Tarutta et al.// Bulletin of Ophthalmology. – 2012. – No. 1.