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SUMMARY

Syringomyelia; pathogenesis, diagnosis, Treatment, and rehabilitation

Syringomyelia is a rare, chronic disease. It can damage the spinal cord due to formation of fluid-filled cysts in central canal. Pathogenesis of idiopathic syringomyelia has not been established so far despite many scientific researches. It seems that the most important is obstruction of cerebrospinal fluid flow in the region of foramen magnum caused by congenital or developmental malformations like Chiari I malformation i.e. cerebellar tonsils herniation into cervical spinal canal. Frequent coexistence of syringomyelia and Chiari I malformation results in overlapping of symptoms caused by these processes.

Because cervical region is the most frequently involved, symptoms first appear in upper extremities. The most characteristic symptoms are: atrophy of small muscles of hands, decreased sense-perception, neuropathic pain, scoliosis and dysfunction of sphincters. When medulla oblongata is involved, speech musculature, phonation and swallowing may be disturbed.

The procedure for identifying syringomyelia for many years was based on the patient's complaints and symptoms, at present the basic examination is magnetic resonance imaging.

Various surgical techniques are used to treat syringomyelia, but they may be divided into three groups: craniocervical decompression, laminectomy with drainage of syringomyelic cyst and ventriculostomy terminal.

Non-surgical therapy includes pharmacological treatment of neuropathic pain and spinal or peripheral electrical stimulation.

The predominant part of rehabilitation is kinetic therapy, assisted by physical and psychological therapy. In the social rehabilitation should be engaged multidisciplinary team consisting of physician, nurse, physiotherapist, psychologist and social worker.

The objective of rehabilitation is to maintain or recover the patient's maximal capabilities, reduce dependence and improve quality of life.