Hydrops Layer with Communicating Hydrocephal- about a Case

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Clinical Case

A 28-year-old female patient presented with a three-month rotational vertigo seizure with concomitant nausea, severe bilateral tinnitus, optic fullness and constant universal pulsating headache.

Personal history

I abort one [1] month before the start of EA
Family background:
No contributarions.

Physical examination: Positive Findings ORL:
Mucosa of pale inferior turbinates with slight hypertrophy.
Bilateral tympanic tumors with Positive Neumatoscopy bilaterally.

Otoneurological study

a) Clinical: Nystagmus positive to extreme gaze, gait test and Romberg Lateralization to the right.
b) Cochlear scan:
c) Tonal audiometry: Normal hearing
d) Vocal Audiometry: Good Bilateral Discrimination.
e) Impedanciometry: Both curves deformation, pressure and normal amplitude.
f) Stapedial reflexes ipsi and contralateral present with evidence of bilateral recruitment.
g) Neural adaptation: Normal.
h) Rinne’s test: positive (+) Bilateral. Weber Test: Indifferent
i) Electronystagmography: Nystagmus spontaneous and semispontaneous absent, Normal tract, Nystagmus Optokinetic present with correct sense
j) Rotatory Chair: presence of both vestibular systems, Caloric Tests: Nystagmus of high frequency to bilateral bithermal irrigation.
k) Otoneurological Conclusion: Normal Hearing. Bilateral labyrinthine irritability.

Laboratory tests

Hematology, Blood Chemistry, Cholesterol, Triglycerides, HIV, VDRL, EX. Urine, within normal limits.

RSM Cerebral

Global Hydrocephalus Communicant with slight Stenosis of the Silvio Aqueduct.
Driving

It is evaluated jointly with Neurosurgery, suggesting Peritoneal Ventricle Derivation, it is intervened surgically obtaining a satisfactory resolution of hydrocephalus with disappearance of the cochleovestibular symptomatology. Currently 3 years postoperative, in control and follow-up, asymptomatic.

Discussion

Labyrinthine Hydrops is defined as the dilatation of labyrinthine membranes associated with an increase in endolaberinic fluid, clinically manifested by Vertigo, Tinnitus, optical Plenitude, and can reach Hearing Loss. Its etiology is unknown. An origin has been postulated:

Embryopathic (Dysplasia of Mondini), Acquired (post infections, trauma, metabolic, endocrine, immunological, vascular disorders), Ideopathic (Meniere’s disease). Mechanical obstruction of the endolymphatic duct and labyrinthine microhomostasis disorders have been considered as pathogenic factors responsible for Labyrinthine Hydrops [1-3]. The constant and balanced composition of the fluids of the inner ear are fundamental for the good functioning of the vestibular cochlea.

Inner ear fluids play important roles

a) Provide an appropriate and specific ionic environment for optimal generation of biopotentials necessary for proper inner ear function.

b) Keep the internal ear pressure balanced; The hydrostatic CSF variations are transmitted by the cochlear aqueduct to the perilymphatic spaces, simultaneously this pressure of the liquid is exerted on the walls of the endolymphatic sac, which causes the pressures to be equal on both sides of the Reissner membrane.

c) They allow the transportation of nutrient principles to the terminal organs and the elimination of metabolic waste products from the cochlea.

d) Medium of transmission of the vibrations that go from the stirrup to the energy transforming centers of the middle ear.

Fluids from the inner ear are in contact with the surrounding tissues, they communicate directly or indirectly with the CSF and / or blood, with a permanent exchange of ions and metabolites, due to the biochemical differences between them [1,4]. Intracranial pressure may modify the intralaberinic pressure directly by the Cochlear Aqueduct or indirectly by the labyrinthine branches of the intracranial vasculature, especially the microvenules [5,6]. The maintenance of microhomostasis of internal ear fluids depends on:

i. Energy-dependent ion pumps.

ii. Constant blood circulation.

iii. LCR-Laberintic Barrier.

There are several situations, conditions, or factors that may disturb maintenance of microhomostasis of endolaberinic fluids, leading to accumulation of ions, metabolites, excess fluid in the inner ear compartments, leading to osmotic imbalance, labyrinthine membrane distention And consequently functional disturbances of balance and hearing [1,7,8]. In the case described above, Global Communicating Hydrocephalus is considered to be the disturbing condition of microhomostasis of endolaberinic fluids, condition of osmotic imbalance, labyrinthine membrane distention and Labyrinthine Hydrops.

Conclusion

In the discussion with Neurosurgery, it is suggested that the mild Stenosis of the Silvio Aqueduct (congenital malformation), until now had been adequately tolerated by the patient, since it allowed good management of cerebral fluid dynamics with adequate CSF flow and drainage, is unknown The cause of increased intracranial water volume (no positive history in the anamnesis), responsible for communicating global hydrocephalus, because there is only limitation in the flow and drainage of the CSF, but no obstruction (Obstructive Hydrocephalus). It could be considered viral or immunological post-infectious etiology that caused fibrosis or adhesions of the arachnoid villi in the sagittal sinus, with limitation in the reabsorption of the CSF. Therefore, taking into account the relationship between CSF and endolaberinic fluids, we consider Hydrocephalus as a causal factor of Labyrinthic Hydrops, in our case, when hydrocephalus was resolved, recovery of microhomostasis was achieved, evidently with the disappearance of the Coclleovestibular symptomatology, therefore we conclude:

a. Endolaberinic fluids directly and indirectly contact the CSF.

b. Hydrocephalus is a disorder of cerebral fluid dynamics that can disturb the microhomostasis of the fluids of the inner ear, conditioning a labyrinthine Hydrops.

c. The RSM Cerebral is a complementary examination in the pathogenesis of the Labyrinthic Hydrops.

References


