HEARING LOSS IN PATIENTS WITH SHUNT-TREATED HYDROCEPHALUS

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ABSTRACT

Hearing loss is a common manifestation of the long-term complications in patients with shunt treated hydrocephalus along with motor development disturbance, cognitive and visual impairment, epilepsy and endocrine disorders.

The aim of the present study was to investigate the alterations of hearing in patients with shunt treated hydrocephalus of non-tumor etiology and at least one year after implantation of ventriculo-peritoneal shunt, as well as their impact on the quality of life of patients.

The study included 70 patients (age range 1.25 years – 21.25 years) with shunted non-tumor hydrocephalus and at least one year after placement of the shunt system. Hearing alterations were proved by measuring the brainstem auditory evoked potentials (BAEP) for children up to 5 years of age and children with mental retardation; audiograms was used for children older than 5 years with normal neuro-psychological development (NPD).

Of the 70 studied patients 17 (24%) had hearing loss (10 bilateral and 7-unilateral) and all of them had sensorineural hearing loss, which is associated with low weight at birth, posthemorrhagic hydrocephalus and brainstem symptoms at the time of diagnosis of hydrocephalus. Hearing pathology was found more often in shunt-treated patients with NPD retardation, poor functional status and low quality of life.

Children with shunt-treated hydrocephalus have hearing loss of sensorineural type. Children with brain stem symptomatology at diagnosing hydrocephalus and children with post-hemorrhagic hydrocephalus show higher risk of hearing loss. Children with shunted hydrocephalus and hearing loss show lower NPD, lower quality of life and lower functional status.

Key words: shunted (shunt-treated) hydrocephalus, hearing loss, quality of life, audiometry, brainstem auditory evoked potentials

REЗЮМЕ

У пациентов с гидроцефалией, прошедших лечение при помощи шунтирующих систем, расстройство слуха, а также нарушения в двигательном развитии, когнитивные, зрительные, появление эпилепсии и эндокринные нарушения являются частыми проявлениями долгосрочных осложнений. Целью настоящей работы является исследование изменений слуха у пациентов с шунтированной гидроцефалией неопухолевого происхождения, которым как минимум за год до этого установлена вентрикуло-перитонеальная шунтирующая система, а также исследование воздействия данных изменений на качество жизни пациентов. Пациенты и методы: Исследование охватывает 70 пациентов (возрастной охват 1.25 – 21.25 лет) с шунтируемой неопухолевой гидроцефалией, которым шунтирующая система установлена как минимум за год до этого. Слуховые изменения были установлены путём измерения слуховых вызванных потенциалов ствола мозга у детей до 5 лет и у детей с умственной отсталостью, аудиометрия была применена в отношении детей старше 5 лет с нормальным нейропсихическим развитием. Результаты: Нейросенсорная потеря слуха была установлена у 17 пациентов (24 %) (у 10 - двусторонняя и у 7 – односторонняя), которая связана у всех с низким весом при рождении, постгеморрагической гидроцефалией и симптомами поражений ствола мозга во время диагностирования заболевания. Слуховая патология была установлена чаще у пациентов, прошедших лечение при помощи шунта, у которых установлены отставание в нейропсихическом развитии, плохое функциональное состояние и ухудшенное качество жизни. Выводы: У детей с гидроцефалией, прошедших лечение при помощи шунта потеря слуха является
INTRODUCTION

The introduction of shunt surgery in the neurosurgical practice in the 60s of the last century helped overcome the complications as a result of increased intracranial pressure (ICP), increase survival and improve quality of life (QOL) in children with shunt-treated hydrocephalus. The better survival of shunt-treated patients allows for analysis of the complex effect of hydrocephalus and its treatment on their cognition, motor function, hearing, vision, occurrence of epilepsy and endocrine disorders.

Hearing loss is a component of the long-term complications observed in shunted patients. Only a few of the available studies have investigated the incidence of hearing loss in patients with implanted ventriculo-peritoneal shunts (VPS). Löppönen et al.\(^1\) reported incidence of 38% of high-frequency sensorineural hearing loss in shunt-treated hydrocephalic children, and Edward et al.\(^2\) found abnormal BAEP in 68.8% of the studied infants with hydrocephalus.

Proceeding from the fact that shunt surgery is associated both with resolution and development of auditory dysfunction, Satzer and Guillaume\(^3\) focused their research on the mechanisms of hearing loss in shunt-treated hydrocephalus. They reported that independent of the indirect mechanisms of hearing loss in shunt-treated patients (mass effect, compromised auditory pathway, complications of premature birth and genetically mediated hydrocephalus with hearing loss), there also exists a direct mechanism (the hydrodynamic theory). According to the latter the very intimate relationship between the cerebrospinal fluid (CSF) and the inner ear fluids permits the origin of relative endo- and perilymphatic hydrops under the conditions of abnormal pressure and abnormal CSF circulation. CSF is associated with the perilymph and the changes in the CSF pressure give rise to alterations of perilymphatic pressure.

Auditory dysfunction may arise in the development of hydrocephalus, as a consequence of shunt operations or as a result of neurosurgical intervention, including puncture and drainage of CSF out of the subdural space. Hearing loss can be categorised into permanent, reversible, early onset, late onset, unilateral and bilateral.

AIM

The aim of our study was to investigate the alterations of hearing loss in patients with shunt-treated hydrocephalus of non-tumor etiology, at least one year after the implantation of VPS, and also find their effect on the quality of life of patients.

MATERIALS AND METHODS

The study included 70 patients with shunt-treated non-tumor hydrocephalus studied at least one year after VPS placement. Their mean age was 6.7 years (1.25 – 21.25 years); 36 of the subjects were males and 34 - females. The following groups of hydrocephalus were found according to the etiology: congenital aqueductal stenosis - 18 patients (25.7%), hydrocephalus with myelomeningocele (MMC) - 8 patients (11.4%), Dandy Walker Syndrome – 5 patients (7.1%), chromosomopathy, craniofacial anomalies and congenital cysts - 5 patients (7.1%), postinfectious hydrocephalus (congenital and acquired) - 6 patients (8.6%) and posthemorrhagic hydrocephalus -28 patients (40%).

Examination of auditory analyzer was performed using brainstem auditory evoked potentials (BAEP) for children up to 5 years and for children with NPD retardation; for children without NPD retardation above the age of 5 years we used audiograms. BAEP testing was done with recording electrodes placed as one active electrode placed on mastoid process (A1/A2) and one reference electrode (Cz). BAEP test was performed using monaural click stimulation 11 c/sec. and the average of 2000 responses. Examination was performed at impedance below 5 kΩ and filters for high frequencies - 3000 Hz, and for low frequencies - 100 Hz. To evaluate BAEP we measured the absolute latency of waves I, III and V and the interpeak intervals I-III, III-V I-V. The examination was performed with patients
in lying position, after eating, during natural sleep or after sedation with chlorpromazine. As an instrument for auditory screening 30 dB nHL is the lowest intensity.

As indicators for abnormal results were accepted increased latencies of wave V or III (or both) and increased interpeak latencies I–V, III–V, with elevated hearing threshold or normal hearing threshold.4

For children under 5 years of age with normal mental development we used tonal threshold audiometry to assess the auditory analyzer. The minimum hearing threshold was defined through the air and bone conduction. For examining air conduction the tones were transmitted with the help of earphones and for bone conduction – with the help of an audiometer vibrator. The patients were examined as they had to respond to definite tones of different frequency and intensity with the aim to define the hearing threshold. The level in dB at which the signal was hardly heard was defined as the minimum hearing threshold. The minimum hearing thresholds were entered into an audiogram. In case of disorder of the sound conducting apparatus of the ear there is sound conductive hearing loss. Hearing loss by air conduction is available. In case of disorder of the sensorineural apparatus (organ of Corti, neurons, sound conduction pathways and central auditory system) there is sensorineural hearing loss. Hearing loss by air and bone conduction is available.

Neuro-psychological development of all patients was examined by a pediatric psychologist. Children up to 3 years of age and those with severe NPD retardation were investigated using Manova-Tomova’s QOD (quotient of development) scale. IQ (Binet-Termann) was tested in children over the age of 3 years. QOD and IQ results were summarized into a summarized NPD. Mental retardation was defined if tested and calculated IQ was below 70, in accordance with the International Classification of Diseases (ICD 10).

The parents of children above the age of 5 years completed the Hydrocephalus Outcome Questionnaire (HOQ) created by Kulkarni et al. It is applicable to patients between 5-18 years of age no matter what the etiology of hydrocephalus is and at least 6 months after making the diagnosis. The questionnaire consists of 51 questions which concern the physical, cognitive and social-emotional health. The highest assessment score is 1.

Because of lack of specific test for QOL assessment in patients with hydrocephalus under 5 years of age, their parents completed the Pediatric Quality of life Questionnaire (PedsQL). It includes physical, emotional, social and cognitive health of healthy children, children with acute diseases and children with chronic diseases. The highest assessment score of the answer is 1. We integrated the results from the investigation of both age categories in a summarized QOL result (total and by domains).

The assessment of the functional status of all patients included in the study was realized using the Functional status scale (FSS) (Pediatrics -2009 American Academy of Pediatrics). It includes examination of mental status, sensorial functioning, communication, motor functioning and respiratory status and feeding. The optimum score is 6, deterioration of functional status is associated with the increase of the digital value.

**Statistical analysis**

The collected primary information was analyzed with SPSS Statistics 19.0, and the diagrams were prepared with Microsoft Office 2010.

The following statistical methods were used: descriptive statistics, non-parametric analysis and graphical analysis. Level of significance for the null hypothesis was accepted at p < 0.05.

**RESULTS**

Of the 70 studied patients with shunted hydrocephalus 17 patients (24.3%) had hearing loss. Ten of these (14.3%) did not hear with both ears, and 7 patients (10%) did not hear with one ear (Fig. 1).

Fifty patients were examined by BAEP testing and 20 were examined using audiometry. We registered auditory sensorineural dysfunction in all patients. Of the 50 patients who were examined by BAEP, 10 patients (20%) did not hear with both ears, 4 (8%) did not hear with the right ear and 1 (2 %) – with the left ear. Of the 20 patients who were examined by audiogram: 1 patient (5%) did not hear with the right ear and 1 (5%) did not hear with the left ear (Fig. 2).

**Relationship between hearing and the rest indicators**

We found no statistically significant difference in gender for hearing loss when examined by BAEP, audiogram or summarized. There is no statistically significant relationship between hearing and gestation age in shunted patients (p > 0.05, χ² = 3.695).

We found statistically significant difference between hearing and weight at birth (p < 0.05, χ² = 5.598). 70.6% of the studied children with weight below 2500 g had hearing loss, while in children
with weight over 2500 g hearing loss was found in 29.4% (Fig. 3).

We found statistically significant difference between hearing and etiology groups of hydrocephalus ($p < 0.05, \chi^2 = 6.155$). The greatest percentage among patients with hearing loss was found in the group with posthemorrhagic hydrocephalus (Fig. 4).

We found significant difference between hearing and bradypnea as a symptom in the diagnosis of hydrocephalus ($p < 0.05, \chi^2 = 5.043$). Patients with manifested bradypnea showed higher incidence of hearing loss when hydrocephalus was diagnosed. More than half of the bradypnea patients had hearing loss when hydrocephalus was diagnosed (Fig. 5).

We did not find statistically significant difference between hearing and delayed placement of a shunting system ($p > 0.05, \chi^2 = 0.186$), as well as between hearing and the number of revisions of the shunting system ($p > 0.05, \chi^2 = 1.916$).

The hearing of our patients with shunted hydrocephalus was not affected by mechanical, functional and infectious complications of the shunting system ($p > 0.05$).
We did not find higher incidence of hearing loss in patients with shunted hydrocephalus and accompanying child cerebral palsy (CCP) \((p > 0.05, \chi^2 = 3.19)\).

We found that the incidence of hearing loss was higher among children with shunted hydrocephalus and with severe NPD retardation \((p < 0.01, \chi^2 = 11.171)\) (Fig. 7).

We found association between the incidence of hearing loss and quality of life both as overall health \((p < 0.05, \chi^2 = 6.371)\) and as physical health alone \((p < 0.001, \chi^2 = 13.704)\) (Figs 8, 9).

We did not find statistically significant difference between hearing loss and the other elements of quality of life.

We found very high statistically significant association between hearing loss and FSS \((p < 0.001, \chi^2 = 22.290)\) – children with high FSS values, correlating with low functional status, showed higher incidence of hearing loss (Fig. 10).

**DISCUSSION**

We found hearing loss in 24% of the patients, a result which is considerably lower in comparison with all reported results in the available literature and is quite similar to the results reported by Löppönen et al.\(^1\) - 38%. We cannot comment upon the reasons for hearing loss in the children from our group. We have no information whether hearing loss occurred before the placement of the shunt or whether there was improvement of hearing after shunting. None of the children had prior examination of hearing loss.

Our data concerning sensorineural hearing loss are consistent with the data reported by Löppönen et al.\(^1\), Sammons et al.\(^7\), Spirakis and Hurley\(^8\), and...
Sandro et al. (1999) on sensorineural hearing loss in patients with shunt-treated hydrocephalus. Our study also confirmed the prevalence of bilateral hearing loss reported by Lopponen et al.1 and Ducati et al.10. We found bilateral hearing deficiency in 58.8% of the children with hearing loss, which was very close to the results of Löppönen et al.1 who found bilateral hearing loss in half of their patients. Our results differ from the results of Spirakis and Hurley8, who reported 100% hearing loss ipsilateral to the inserted CNS catheter – for this reason we cannot confirm the association between hearing loss and the presence of a shunt catheter in spite of the higher percentage of hearing loss ipsilateral of the catheter in our patients.

We corroborated the role of etiology of hydrocephalus as a risk factor for hearing loss which was reported by Kraus et al.11 and Ducati et al.10. We found normal hearing most often in aqueductal stenosis which was consistent with the data reported by Ducati et al.10. The results of our study showed the highest frequency of hearing loss in children with post-hemorrhagic hydrocephalus.

Our results showed higher incidence of hearing loss in patients who, when hydrocephalus was diagnosed, had bradypnea and abnormal respiratory rhythm as manifestations of brain stem symptomatology. So we confirmed the thesis of Kraus et al.11 on the importance of brain stem symptomatology as a risk factor for hearing loss in shunt-treated patients.

We also confirm the causal relationship between hearing loss and abnormal NPD, lower quality of life and lower functional status.

CONCLUSIONS
1. Children with shunt-treated hydrocephalus have hearing loss of sensorineural type.
2. Children with shunt-treated hydrocephalus and brain stem symptomatology when diagnosing hydrocephalus, as well as children with post-hemorrhagic hydrocephalus show higher risk of hearing loss.
3. Children with shunt-treated hydrocephalus and hearing loss show lower NPD, lower quality of life and lower functional status.

REFERENCES
7. Sammons VJ, Jacobson E, Lawson J. Resolution


