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Audiologic evaluations of children with mucopolysaccharidosis

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KEYWORDS
Mucopolysaccharidosis;
Hearing loss;
Audiologic evaluation

Abstract
Introduction: Mucopolysaccharidosis is a hereditary lysosomal storage disease, which develops due to a deficiency in the enzymes that play a role in the metabolism of glycosaminoglycans (GAG). The incidence of mucopolysaccharidosis is 1/25,000, with autosomal recessive inheritance (except for MPS II). Mucopolysaccharidosis occurs in seven different types, each with a different congenital deficiency of lysosomal enzymes. In mucopolysaccharidosis patients, even though progression of clinical findings is not prominent, the disease advances and causes death at early ages. Facial dysmorphism, growth retardation, mental retardation, and skeletal or joint dysplasia are the most frequently found symptoms in these patients.
Objective: The purpose of our study is to present the types of hearing loss types and tympanometric findings of patients with mucopolysaccharidosis referred to our clinic with suspicion of hearing loss.
Methods: After otorhinolaryngological examination, 9 patients with different types of mucopolysaccharidosis, underwent to immittance and audiometric evaluations, performed according to their physical and mental abilities, and ages, in order to determine their hearing thresholds.
Results: The audiometric findings of the 9 patients followed with mucopolysaccharidosis were reported separately for each case.

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Conclusion: Based on the high frequency of hearing loss in mucopolysaccharidosis patients, early and detailed audiological evaluations are highly desirable. Therefore, regular and systematic multidisciplinary evaluations are very important.
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Introduction

Mucopolysaccharidosis (MPS) is a hereditary lysosomal storage disease, which develops due to the deficiency in the enzymes that play a role in the metabolism of glycosaminoglycans (GAG). Progressive GAG accumulation causes advanced growth retardation, skeletal deformities, hearing loss, weak joint movement, and facial dysmorphism. In accordance with the deficiency of the known 11 enzymes, MPS has seven sub-types (MPS I, MPS II, MPS III, MPS IV, MPS VI, MPS VII, MPS IX). Although the incidence of MPS in accordance with its sub-types varies, a general incidence of 1/25,000 has been reported.1-3

In patients with MPS, hearing loss is observed due to various reasons. Conductive hearing loss (CHL) may develop due to recurrent upper respiratory tract infections and serous otitis media or bone chain deformities. Sensorineural hearing loss (SHL) is thought to be caused due to the accumulation of GAG in the cochlea, auditory nerve, and brain stem. In many patients, mixed-type hearing loss, along with the symptoms of conductive and sensorineural hearing loss, can also be seen. In CHL related to middle ear effusion, while the ventilation tube implementation is frequently used as treatment, hearing aids are advised in SHL.4-7 Early diagnosis of hearing losses carries great importance in terms of early intervention in MPS patients, as well as in all other patients.8,9

The purpose of the present study was to present the hearing loss type, degree, and tympanometric findings of patients with MPS, who are directed to this clinic with the suspicion of hearing loss.

Methods

The nine patients who had been directed to this department with the suspicion of hearing loss, whose follow-up had been conducted after they have been diagnosed with MPS, were included in this study (Ethical Committee approval number: GÜKAEK-461). The otological examinations of the patients were performed by the otorhinolaryngologist. Information related to each patient’s otorhinolaryngological complaints and treatments were compiled and middle ear infection, ear discharge, tinnitus, dizziness, tympanum problems, adenoidectomy, ear tube placement, and hearing aid usage were determined. After the otorhinolaryngological examination, the immitancemetric evaluations were conducted and...
Table 1  Degree of hearing loss (dB HL).

<table>
<thead>
<tr>
<th>Degree of hearing loss</th>
<th>Hearing loss range (dB HL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td>−10 to 15</td>
</tr>
<tr>
<td>Slight</td>
<td>16−25</td>
</tr>
<tr>
<td>Mild</td>
<td>26−40</td>
</tr>
<tr>
<td>Moderate</td>
<td>41−55</td>
</tr>
<tr>
<td>Moderately severe</td>
<td>56−70</td>
</tr>
<tr>
<td>Severe</td>
<td>71−90</td>
</tr>
<tr>
<td>Profound</td>
<td>90+</td>
</tr>
</tbody>
</table>

Table 2  Tympanogram types (daPa).

<table>
<thead>
<tr>
<th>Tympanogram type</th>
<th>Pressure intervals (daPa)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Type A</td>
<td>+50 to (−150)</td>
</tr>
<tr>
<td>Type B</td>
<td>−400</td>
</tr>
<tr>
<td>Type C</td>
<td>−150 to (−399)</td>
</tr>
</tbody>
</table>

Statistical analysis

For the statistical analysis, SPSS version 16.0 (SPSS Inc., Chicago, IL, United States) was used. For the age average and standard deviation (SD), descriptive statistics were developed.

Results

The age average of the total of nine patients with MPS, four female (44.44%) and five male (55.55%), was determined as 5.66 ± 2.73. There were three cases of MPS VI (33.33%), three cases of MPS I (33.33%), one case of MPS III (11.11%), and two cases of MPS IV (22.22%) that were diagnosed. Cases MÖ and EÖ are siblings; İK and İK 2 are twins. The other five patients are not related. The degree of hearing loss, its type, and tympanometrical findings are shown in Table 3.

Cases MÖ and EÖ were concordant, with hearing thresholds and air-bone gaps being equal (daPa). There were six ears with pure conductive hearing loss (33.33%) (HL one ear: slight; HL two ears: mild; HL four ears: moderate, and HL two ears: moderately severe); 12 ears with mixed hearing loss (66.66%) (HL two ears: mild; HL four ears: moderate, HL five ears: severe, and HL one ear: profound) were determined. There were 14 ears with Type B tympanogram (77.77%), three ears with Type C tympanogram (16.66%), and one ear with Type A tympanogram (5.55%).

E.C., B.E.B., and MÖ had one tympanogram. Although the other patients were advised regarding the VT application as well, it has not been carried out upon the wish of their families. With the exception of MÖ and EÖ, although all patients were advised to use hearing aids, the families did not adopt the use of these devices and did not come regularly for follow-ups.

Discussion

In cases with MPS (Table 4), the initiation of treatment in early stages carries great importance in terms of the prognosis of the disease. However, due to the late appearance of the symptoms, the definite diagnosis range is 3–4 years of age. In the follow-up of the stages of the treatments of patients with MPS, the otolaryngologist and audiologists have a significant role in multidisciplinary teamwork. Recurrent serous otitis media and hearing loss
Table 4 Air and bone conduction thresholds of patients with MPS (dB HL).

<table>
<thead>
<tr>
<th>Patient</th>
<th>Average air conduction thresholds (500-4000 Hz)</th>
<th>Average bone conduction thresholds (500-4000 Hz)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Right</td>
<td>Left</td>
</tr>
<tr>
<td>MÖ</td>
<td>22.5</td>
<td>26.25</td>
</tr>
<tr>
<td>EÖ</td>
<td>65</td>
<td>57.5</td>
</tr>
<tr>
<td>İk1</td>
<td>70*</td>
<td></td>
</tr>
<tr>
<td>İk2</td>
<td>70*</td>
<td></td>
</tr>
<tr>
<td>EÇ</td>
<td>46.25</td>
<td>60</td>
</tr>
<tr>
<td>SD</td>
<td>47.5</td>
<td>38.75</td>
</tr>
<tr>
<td>BEB</td>
<td>51.25</td>
<td>56.25</td>
</tr>
<tr>
<td>MSE</td>
<td>43.75</td>
<td>45</td>
</tr>
<tr>
<td>YCE</td>
<td>81.25</td>
<td>90</td>
</tr>
</tbody>
</table>

* The air conduction thresholds have been determined in free field.
+ The bone conduction thresholds have been determined in accordance with the speech recognition thresholds.

that develops in individuals within this disease group are the reasons why better, long-term follow-ups should be done. Hearing loss can be seen in various types and degrees in MPS in general. Overall, conductive hearing losses due to chronic effusion and eustachian tube dysfunction are seen more in patients with MPS; the incidence of SHL has not been determined very clearly.10

Although conductive and mixed-type hearing loss were observed in the cases of the present study, SHL compatible with literature has not been determined. In the studies conducted, it was stated that the incidence of mixed hearing loss is more, and that this fact is related with age. While conductive hearing loss is determined in early ages and in patients who are diagnosed earlier, it has been stated that mixed hearing loss can be observed, and that this rate is between 33% and 71% as age increases. In the present cases, degree of hearing loss was determined from slight to severe. This difference is considered to be due to GAG’s excessive accumulation in the cochlear ducts, stria vascularis, and cochlear nerve, which prevents them from functioning efficiently.14 In the present cases, the incidence of Type B and C tympanogram was determined to be high, in accordance with the literature. According to Yu Lin et al., while this rate is stated to reach 70%, the incidence of Type B tympanogram was determined as 77.77% and the incidence of Type C tympanogram was determined as 16.66% in the present study, similarly. VT application is advised to cases with similar clinical findings in different clinics as well, just as in the present study. However, a majority of the cases in this study have hindered their treatments, due to their families’ rejection of the VT application. In cases for which hearing aids are advised, while socio-economic inadequacy creates hardships in coming to follow-ups and obtaining these aids, the fact that many live in different cities prevents continuity in follow-ups.

Conclusion

In cases with MPS, since the incidence of hearing loss is high, the hearing loss should be determined in early stages. After the diagnosis, the required treatments should be started promptly and thus quality of life should be increased. However, it is considered that this is only possible as a result of regular and systematic follow-ups and with multidisciplinary studies.

Conflicts of interest

The authors declare no conflicts of interest.

References