Effects of Obesity on the Auditory Function of Children and Adolescents

Engin Başer1 Havva Nur Peltek Kendirci2

1Department of Otorhinolaryngology, Head and Neck Surgery, University of Health Sciences, Izmir Tepecik Training and Research Hospital, Izmir, Turkey
2Department of Pediatric Endocrinology, Hitit University, Faculty of Medicine, Corum, Turkey


Abstract

Introduction  Childhood and adolescent obesity is associated with insulin resistance, abnormal glucose metabolism, hypertension, dyslipidemia, inflammation, liver disease, and compromised vascular function.

Objective  We aimed to evaluate the effects of obesity on the auditory function and speech audiometry of children and adolescents.

Methods  Subjects with a body mass index (BMI) higher than +2 standard deviation (SD) were classified as obese, and subjects with normal BMI SD were classified as the control group. Blood samples were taken for glucose, insulin, and lipid profiles following an 8-hour fasting period, and a hepatobiliary ultrasound was performed. The homeostatic model assessment for insulin resistance (HOMA-IR) was calculated. The audiological evaluation included pure-tone audiometry (PTA), speech reception threshold (SRT), and speech discrimination score (SDS).

Results  The study included 100 children (50 girls) with obesity, with a mean age of 11.4 ± 2.9 years and 30 children with normal body weight, with a mean age of 11.9 ± 3.3 years. Of the children with obesity, 55% (n = 55) were found to have hyperlipidemia, 68% (n = 68) insulin resistance, and 21% (n = 21) hepatosteatosis. There were no statistically significant differences between children with obesity and the control group in terms of SDS or PTA, while SRT was found to be higher in children with obesity. There was no difference between obese children with or without hyperlipidemia, between obese children with or without insulin resistance, and between obese children with or without hepatosteatosis, according to hearing tests.

Conclusion  The result of the present study indicates that children with obesity are more prone to having auditory problems than the normal population. We recommend more frequent audiological evaluations, including speech audiometry, in children and adolescents with obesity problems.

Keywords  ➤ childhood  ➤ adolescent  ➤ obesity  ➤ pure-tone audiometry  ➤ speech audiometry

received September 2, 2020  accepted after revision March 23, 2021  published online January 24, 2022


© 2022. Fundação Otorrinolaringologia. All rights reserved. This is an open access article published by Thieme under the terms of the Creative Commons Attribution-NonDerivative-NonCommercial-License, permitting copying and reproduction so long as the original work is given appropriate credit. Contents may not be used for commercial purposes, or adapted, remixed, transformed or built upon. (https://creativecommons.org/licenses/by-nc-nd/4.0/)

Thieme Revinter Publicações Ltda., Rua do Matoso 170, Rio de Janeiro, RJ, CEP 20270-135, Brazil
Introduction

Pediatric obesity is one of the most serious global health problems of this century. Obesity occurring in childhood may lead to several complex metabolic complications, including increased insulin resistance, diabetes, respiratory problems, musculoskeletal system disorders, and predisposition to certain forms of cancer, as well as cardiovascular complications, and the psychological effects of low self-esteem, low academic achievement, and depression. In addition, it is reported that obesity starting in childhood is more harmful than obesity starting in adulthood.3

In recent years, the relationship between obesity and hearing loss has attracted the attention of researchers. The cochlear hair cells of the inner ear are fundamental receptors in hearing function. Mature mammalian cochlear hair cells are unable to regenerate. Therefore, damage to these cells results in permanent hearing loss.2 In a few studies performed through microvascular circulation mechanisms of the cochlea, obesity was defined as a predisposing factor for hearing loss.4 It is known that obesity causes dysfunction in many organs and sensorial systems of the body via oxidative stress and lipotoxicity.5 Many comorbid conditions, such as metabolic, cardiovascular, orthopedic, neurological, hepatic, pulmonary, and renal disorders, have been associated with childhood obesity. Until recently, many of the aforementioned health conditions had only been observed in adults, whereas now they are extremely prevalent in obese children. Childhood obesity is also associated with poor academic performance and the child experiencing lower quality of life.6 Since individuals with hearing loss also experience communication loss, in addition to other obesity-related comorbidities, this condition may increase obesity much more by leading to social regression. Obesity and hearing loss considerably reduce both physical and psychological quality of life.7

We believe that the effect of obesity and its comorbidities starting in childhood, which is an important stage of cognitive development, on hearing has not been examined as much as it deserves in the literature. We aimed to evaluate the effects of obesity on the auditory function and speech audiometry of children and adolescents as well as to investigate the relationships between audiological parameters and comorbidities such as insulin resistance, dyslipidemia, and hepatosteatosis in obese children.

Methods

The study protocol was approved by the local ethics committee (Approval number: 108400987–281). Written informed consent was obtained from the parents at the start of the study in accordance with the Declaration of Helsinki.

Study Procedures

A total of 100 obese children and adolescents between the ages of 5 and 17 with a BMI > 95% who presented to the pediatric endocrinology outpatient clinic between May and October 2014 and who were cooperative when undertaking audiometric examination were included in the study. Thirty children and adolescents with a normal body weight who underwent audiometry for school screening and were age-gender matched with the patient group constituted the control group. The children and adolescents with a type A tympanogram results and normal otorhinolaryngological examination results, and without comorbid systemic disease, without history of use of ototoxic medications, without acoustic and/or physical trauma, or without history of hearing loss in the family were included in the study. Children and adolescents with syndromic obesity, known hearing disorder, familial hearing loss, upper respiratory tract infection within the last 3 months, prominent hypertrophy of adenoid tissue with or without tonsil enlargement, and tympanic membrane perforation or cicatricial stricture were excluded from the study.

Following overnight fasting, venous blood samples were collected for the measurement of serum levels of triglycerides, cholesterol, insulin, and glucose. Insulin resistance was assessed with the homeostasis model assessment of insulin resistance (HOMA-IR), which was calculated by using the following formula: [fasting insulin (µU/L) x fasting glucose level (mmol/L)/22.5]. Liver ultrasound was performed in all patients. A value of HOMA-IR > 2.5 in prepubertal children and > 3.1 in pubertal children and adolescents was defined as insulin resistance.8 The definition of dyslipidemia was made according to the triglyceride and high-density lipoprotein (HDL)-cholesterol levels in Table 1.9

Hearing Evaluation

Audiometry tests were performed in a sound-treated room and at frequencies of 125–8,000 Hz. Pure tone average (PTA), speech reception threshold (SRT), and speech discrimination scores (SDS) were measured at frequencies of 250, 500, 1,000, 2,000, 4,000, and 8,000 Hz. Pure tone average was calculated by averaging measurements at 500, 1,000, 2,000, and 4,000 Hz. Speech discrimination score was evaluated with monosyllabic words. Audiometric evaluations were

Table 1 The definition of dyslipidemia

<table>
<thead>
<tr>
<th>HDL-cholesterol (mg/dl)</th>
<th>5–9 years old</th>
<th>10–14 years old</th>
<th>15–18 years old</th>
</tr>
</thead>
<tbody>
<tr>
<td>Boy</td>
<td>Girl</td>
<td>Boy</td>
<td>Girl</td>
</tr>
<tr>
<td>&lt; 42</td>
<td>&lt; 38</td>
<td>&lt; 40</td>
<td>&lt; 40</td>
</tr>
<tr>
<td>Triglyceride (mg/dl)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&gt; 85</td>
<td>&gt; 126</td>
<td>&gt; 115</td>
<td>&gt; 120</td>
</tr>
</tbody>
</table>

Abbreviation: HDL, high-density lipoprotein.
performed by a single senior audiometrist. Physical examinations of the patients were performed by the same otolaryngologist and pediatric endocrinologist.

**Statistical Analysis**

The IBM SPSS Statistics for Windows Version 22.0 software (IBM Corp., Armonk, NY, USA) was used for statistical analysis. Descriptive statistics of the variables studied were presented as mean and standard deviation (SD) for continuous variables. Descriptive statistics of categorical variables were expressed as counts and percentages. Skewness and kurtosis tests were used to determine whether or not data were normally distributed. The Student t-test was used for intergroup comparisons of variables with normal distribution. A p-value of < 0.05 was considered to be statistically significant.

**Results**

The present study included 100 obese children (50 girls) with a mean age of 11.4 ± 2.9 years, and 30 children (15 girls) with a normal body weight and a mean age of 11.9 ± 3.3 years. Seventy-five percent (n = 75) of obese subjects and 66.7% (n = 20) of non-obese subjects were determined to be within the pubertal period. No difference was determined between groups regarding age, gender distribution, height, height SDS, and pubertal stage. The clinical characteristics of groups are shown in **Table 2**.

It was observed that 55% (n = 55), 68% (n = 68), and 21% (n = 21) of obese children were determined to have hyperlipidemia, insulin resistance, and hepatosteatosis, respectively. The laboratory results of obese children are shown in **Table 3**.

No statistical difference was determined between obese female children (n = 50) and obese male children (n = 50) in terms of age, weight, weight SDS, height, height standard deviation (SD), BMI, and BMI SD. It was determined that 10% of female children were in the prepubertal period, while 40% of male children were in the prepubertal period and this difference was statistically significant (p = 0.001). There was no difference between female and male children in regards to the presence of hyperlipidemia, insulin resistance, and hepatosteatosis. No difference was determined between female and male children regarding any of the hearing tests (p > 0.05).

No difference was determined between groups with and without hyperlipidemia, and with and without insulin resistance regarding hearing tests (p > 0.05).

The prevalence of hyperlipidemia and hyperinsulinemia was found to be higher in the hepatosteatosis group (p = 0.018, p = 0.021, respectively). While there was no difference between obese subjects with and without hepatosteatosis according to PTA and SDS values (p = 0.399, p = 0.555, respectively), SRT values were determined to be significantly higher in the hepatosteatosis group (p = 0.046).

There was no difference between obese children and control groups in terms of SDS, and PTA, while SRT was found to be higher in children with obesity (**Table 4**).

**Discussion**

Obesity and concomitant comorbid disorders can affect many sensory systems and organs. Pediatric screenings of hearing to identify possible hearing loss and improve loss of quality of life are conducted in school-aged children up to adolescence in many developed countries. PTA, today’s standard in the assessment of hearing, is unsuitable for evaluating its most important attribute of hearing, its cognitive aspect.10 Speech discrimination scores and SRT are more important, considering the development of speech and language, especially in a noisy classroom environment. It may be important to evaluate the effect of obesity, which can be accompanied by many comorbid diseases, on hearing functions in the pediatric-adolescent age group. Therefore, we

| Table 2 Clinical characteristics of the subjects (mean ± SD) (min–max) (mean) |
|-----------------|-----------------|-----------------|-----------------|-----------------|
|                 | Obese group (n = 100) | Control group (n = 30) | P               |
| Age (years)     | 11.4 ± 2.9 (5.4–17.7) | 11.9 ± 3.3 (6.3–17.9) | 0.38            |
| Weight (kg)     | 66.2 ± 19.2 (29.0–113.8) | 43.0 ± 15.9 (18–69) | 0.00            |
| Weight SDS      | 2.7 ± 0.7 (1.0–5.3) | –0.2 ± 0.9 (–1.9–1.6) | 0.00            |
| Height (cm)     | 149.2 ± 14.0 (117.5–183.6) | 150.0 ± 18.6 (110.0–177.2) | 0.70            |
| Height SDS      | 0.64 ± 1.29 (–2.9–4.9) | 0.21 ± 0.83 (–2.0–1.66) | 0.09            |
| BMI (kg/m²)     | 29.0 ± 4.1 (20.4–45.8) | 18.1 ± 3.2 (12.9–24.0) | 0.00            |
| BMI SDS         | 2.6 ± 0.4 (2.0–4.0) | –0.39 ± 1.0 (–2.2–1.4) | 0.00            |
| Puberty (mean)  | 3 ± 5 | 3 ± 5 | 0.97 |

| Table 3 Laboratory features of obese children |
|-----------------|-----------------|-----------------|-----------------|
| Cholesterol (mg/dL) | 176.8 ± 31.1 | 106.0 | 243.0 |
| Triglyceride (mg/dL) | 119.2 ± 62.2 | 34 | 385 |
| HDL (mg/dL)      | 41.7 ± 10.1 | 21 | 68 |
| LDL (mg/dL)      | 100.9 ± 23.7 | 56 | 151 |
| VLDL (mg/dL)     | 33.9 ± 11.5 | 5 | 64 |
| HOMA-IR          | 4.7 ± 2.8 | 1.6 | 14.4 |

Abbreviations: BMI, body mass index; cm, centimeters; kg, kilograms; n, number; SD, standard deviation; SDS, speech discrimination score.
Table 4 Hearing tests of obese subjects and control group (mean ± SD) (min–max)

<table>
<thead>
<tr>
<th></th>
<th>Obese group (n = 100)</th>
<th>Control group (n = 30)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Right</td>
<td>Left</td>
<td></td>
</tr>
<tr>
<td>PTA</td>
<td>7.9 ± 3.5 (2–21)</td>
<td>7.6 ± 3.5 (4.8–19.8)</td>
<td>0.51</td>
</tr>
<tr>
<td></td>
<td>7.7 ± 4.0 (3–28)</td>
<td>7.7 ± 3.5 (4.8–25.0)</td>
<td>0.95</td>
</tr>
<tr>
<td>SRT</td>
<td>17.1 ± 4.9 (10–35)</td>
<td>14.0 ± 4.1 (5.0–25.0)</td>
<td>0.002</td>
</tr>
<tr>
<td></td>
<td>17.0 ± 5.0 (10–35)</td>
<td>14.5 ± 5.0 (5.0–30.0)</td>
<td>0.01</td>
</tr>
<tr>
<td>SDS</td>
<td>90.9 ± 6.6 (76–100)</td>
<td>92.2 ± 5.6 (80.0–100.0)</td>
<td>0.26</td>
</tr>
<tr>
<td></td>
<td>91.7 ± 5.5 (76–100)</td>
<td>92.1 ± 5.1 (84.0–100.0)</td>
<td>0.18</td>
</tr>
</tbody>
</table>

Abbreviations: PTA, pure tone average; SDS, speech discrimination score; SRT, speech reception threshold.

aimed to evaluate the effect of childhood obesity on cochlear functions and neurological pathways by using PTA, SDS, and SRT.

In studies investigating the effects of obesity on the hearing function, results varied according to age groups. In a study evaluating adults, a positive relationship was observed between high BMI and hearing thresholds and a similar effect was seen at all frequencies. Lalwani et al. reported that obesity in adolescents aged 12 to 19 years was associated with sensorineural hearing loss (SNHL) at low frequencies; they found that obesity was associated with a 1.85 fold increase in SNHL, with age and gender being irrelevant. Moore et al. reported that PTA did not vary substantially depending on cognitive ability, but SRT depended strongly on cognitive performance. We determined that PTA was not affected at all frequencies. However, SRT was higher (p < 0.05) and SDS was lower (p > 0.05) in obese subjects. Considering these high SRT and low SDS values, we believe that obese children and adolescents may have a higher risk of developmental delay in speech and language, leading to learning problems. For this reason, SDS and SRT should be added to follow-up of obese children.

Many studies have observed that the relationship between obesity and hearing loss was stronger in females, when compared with males. Shargorodsky et al. concluded that obesity was not a risk factor for hearing loss in male subjects aged 40 to 74 years. Kim et al. determined that there was a positive relationship between PTA and visceral adipose tissue in adult women. Hwang et al. reported that obesity was associated with hearing loss for high frequencies in female subjects older than 55 and for both low and high frequencies in males younger than 55. Similarly, Ucler et al. also concluded that auditory thresholds were elevated at high frequencies in obese adult females. In our study, no difference was determined between female and male subjects, according to hearing tests. This implies that gender-related difference in the association between obesity and hearing functions becomes more pronounced with increasing age.

Although different hypotheses have been suggested regarding the relationship between obesity and hearing loss, the main cause is vasoconstriction in the inner ear. Since the cochlea is a very metabolically active organ, its vascularization and oxygenation are important. Hwank et al. demonstrated that hearing threshold levels at high frequencies were elevated in mice with diet-induced obesity. In the histological study, constricted blood vessels were observed in the stria vascularis part of the cochlea. Moreover, high BMI and dyslipidemia lead to narrowing of arteries and decreased blood flow by causing atherosclerotic vascular disease. Adipose tissue affects insulin resistance, energy metabolism, and atherosclerosis through the release of hormones and cytokines, and it may exacerbate inflammation and end-organ damage caused by obesity. Obesity directly leads to worsening of hearing through lipotoxicity and related oxidative stress or indirectly contributes to peripheral hearing degeneration via its comorbidity-related angiopathy and/or neuropathy.

Satar et al. reported that dyslipidemia could cause cochlear injury by leading to edema in the outer hair cells and the stria vascularis. Lee et al. emphasized that elevated total cholesterol and triglyceride (TG) levels, and increased BMI were associated with increased prevalence of SNHL, which occurred as a result of damage due to cochlear ischemia. Shargorodsky et al. reported that hypercholesterolemia was associated with increased risk of hearing loss in adult males. Evans et al. reported that chronic dyslipidemia associated with elevated TG levels may reduce hearing function. In the study performed by Frederiksen et al., while a strong association was determined between high TG, high BMI, and high HDL levels and hearing loss, there was no association between low-density lipoprotein (LDL) levels and hearing loss. Nevertheless, there are also studies that report a weak association between serum lipid levels and hearing loss. We determined no difference between children/adolescents with and without hyperlipidemia regarding hearing tests. The fact that hyperlipidemia-related auditory function is unaffected in the young population suggests that the duration of exposure may affect auditory function as much as hyperlipidemia itself.

Zivkovic-Marinkov et al. conducted a study on hearing loss in adult patients with type 2 diabetes and reported that patients with type 2 diabetes had significantly impaired hearing compared with the control group, also suggesting that prolongation of poorly-controlled glycemia exacerbated hearing loss. Kilic et al. investigated the effects of metabolic syndrome on hearing functions, and observed that children with metabolic syndrome had higher hearing thresholds, especially at low frequencies. To the best of our knowledge, there is no study investigating the effect of isolated insulin resistance on auditory functions. In our study, we determined no difference between obese children/adolescents with and without insulin resistance regarding hearing tests. Since it does not impair glycemic control, insulin resistance may not affect hearing tests.
Healthy hearing depends on sensitive ears and adequate brain processing. The relationship between obesity and increase in speech threshold may be due to impaired cognitive functions. Childhood obesity is related to decrease in executive function, attention, mental rotation, mathematics, and reading achievement. Obese adolescents have deficits in a range of cognitive functions, such as attention and executive functions. Impairment of audiological parameters, as mentioned by many studies, may be due to decrease in cognitive function besides the microvascular circulatory disorder at the cochlear level. This should be exposed with large and comprehensive studies.

Language skills are also important determinants of daily functioning and health and are closely linked to academic and employment outcomes. Speech perception (measured as SRT) develops rapidly toward adolescence and then more slowly until it plateaus in mid-adulthood, before declining at ~60 years of age. Changes in hearing during childhood and adolescence can be neglected by the families and physicians, and this condition may adversely affect the social development, academic performance, and cognitive functions of children. It should be kept in mind that negative social and cognitive outcomes resulting from hearing loss may exacerbate the clinical manifestation of obesity in individuals even further. The struggle against obesity, especially in childhood, is critical to the social and intellectual health of society. According to our results, we believe that hearing evaluation with speech audiometry is necessary, regardless of the etiology of hearing impairment, such as microvascular circulation imbalance, psychosocial effects, and cognitive dysfunction. Moreover, audiologic follow-up should be conducted more frequently than in the normal childhood hearing screening program.

The present study has several limitations. First, the number of patients in our group was not large. Secondly, our study had a cross-sectional design. For this reason, we were unable to evaluate the auditory functions and speech audiometry of our subjects in the long term. The tests were also subjective. Therefore, additional long-term prospective studies with a greater number of patient and control groups are required.

Conclusion

Obesity in childhood is associated with higher SRT and lower SDS. We suggest that audiological functions (with speech audiometry) should be followed up more frequently in obese children and adolescents.

Informed Consent

“Informed consent was obtained from all individual participants included in the study.” Written consent was taken from the parents at the beginning of the study, in accordance with the Declaration of Helsinki.

Financial Disclosure

No financial disclosure.

Conflict of Interests

The authors declare that they have no conflict of interests.

References


