Research Article

The Importance of Saccular Function to Motor Development in Children with Hearing Impairments

Mary S. Shall

Department of Physical Therapy, Virginia Commonwealth University, P.O. Box 980224, Richmond, VA 23298-0224, USA

Correspondence should be addressed to Mary S. Shall, msshall@vcu.edu

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Children with hearing deficits frequently have delayed motor development. The purpose of this study was to evaluate saccular function in children with hearing impairments using the Vestibular Evoked Myogenic Potential (VEMP). The impact of the saccular hypofunction on the timely maturation of normal balance strategies was examined using the Movement Assessment Battery for Children (Movement ABC). Thirty-three children with bilateral severe/profound hearing impairment between 4 and 7 years of age were recruited from a three-state area. Approximately half of the sample had one or bilateral cochlear implants, one used bilateral hearing aids, and the rest used no amplification. Parents reported whether the hearing impairment was diagnosed within the first year or after 2 years of age. No VEMP was evoked in two thirds of the hearing impaired (HI) children in response to the bone-conducted stimulus. Children who were reportedly hearing impaired since birth had significantly poorer scores when tested with the Movement ABC.

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1. Introduction

Most newborn infants are screened for hearing impairment prior to leaving the hospital. As a result, most parents are prepared for the choices of communication needs during this most important period of development. On the other hand, little attention is paid to the performance of the vestibular sensory system, whose receptors are located in the same area as the cochlear receptors. The vestibular nuclei provide a consistent input to the postural muscle motoneurons particularly during initial postnatal development [1]. The lateral vestibulospinal tract affects the vestibulospinal reflex by synapsing directly with motoneurons innervating extensor muscles and indirectly through spinal cord interneurons [2]. Obviously, the development of righting and locomotor skills should parallel sensory system development and motivation. This sensorimotor development has been more extensively studied in nonhuman mammals as they develop motor skills during the first four months [3, 4]. The ferret is born with sensory systems that are more immature than that of the rat or cat. The eyes open at about postnatal day 28 (P28) [5] and hearing becomes functional at about P32 [6]. The vestibular receptors mature at about the same rate as the auditory system; the maculae of the saccule and utricle mature first (~P10) and the cristae of the semicircular canals develop a little more slowly to reach the adult form at about 24 days after birth [7], and the ferrets begin walking at about P24. Neonatal removal of the vestibular labyrinths in ferrets [8] and rats [9] has resulted in changes in muscle physiology and muscle fiber type, persistent unstable standing balance, and reduced velocity of locomotion. Animals that undergo labyrinthectomies as a juvenile or an adult may have some loss of balance reactions but adapt quickly and do not change the anatomy and physiology of the muscles.

In contrast, vestibular receptors have been seen under the microscope in 7–9-week-old human embryos [10], but the vestibular reflexes develop a little more slowly in humans as the sensory systems continue to mature and integrate. Shumway-Cook and Woollacott proposed that, normally, children undergo transition to more adult strategies of posture control between the ages of 4 and 6 [11]. Rine et al. [12] examined the gross motor development among young children with bilateral moderate to profound sensorineural hearing impairment (HI). Delayed gross motor development was evident in these children and a majority received “vestibular hypofunction scores” using the Southern
implants for at least one year prior to testing (range bilateral hearing aids. Nineteen of the children had cochlear 2 were 4 years old and 1 was 5 years old. One child had refused the test either prior to any testing or during the test; 66.1 = All children were between the ages of 4 and 7 years (mean gene but most had not been tested for genetic markers. other children had tested positively for the connexin 26 they had a cytomegalovirus infection during pregnancy. Two unknown in the majority of the cases. Two mothers said Carolina Triad Hitch-Up. The cause of hearing loss was with hearing impairment from birth with a group of children that lost their hearing after 2 years of age.

2. Methods

Thirty-three children (20 males, 13 females) with bilateral severe/profound sensorineural hearing impairments (HIs) (loss greater than 71 dB) were recruited as a sample of convenience through parent support groups such as the Virginia Chapter of the AG Bell Association, the Speech-Language-Hearing Association of Virginia, and the North Carolina Triad Hitch-Up. The cause of hearing loss was unknown in the majority of the cases. Two mothers said they had a cytomegalovirus infection during pregnancy. Two other children had tested positively for the connexin 26 gene but most had not been tested for genetic markers. All children were between the ages of 4 and 7 years (mean = 66.1 ± 12.3 months). There were 3 other children that refused the test either prior to any testing or during the test; 2 were 4 years old and 1 was 5 years old. One child had bilateral hearing aids. Nineteen of the children had cochlear implants for at least one year prior to testing (range = 1– 5 years). Five of the nineteen children only had a cochlear implant in one ear. Parents or guardians completed forms providing a brief health history on the child and consented to perform the evaluation. Most presented audiological documentation. Twenty-four of the children had hearing impairment since birth but no other obvious neurologic damage.

The Vestibular Evoked Myogenic Potential (VEMP) was used to evaluate saccular function. The child lay supine on a mat on the floor with the investigator in a separate quiet room and frequently a parent was in the immediate proximity. The skin over the sternocleidomastoid (SCM) muscles and sternum were cleaned with an alcohol wipe. A surface bar electrode (2 concave Ag/AgCl electrodes separated by 3 cm and coated with conductivity gel) was placed on the SCM muscle (active on the muscle belly, reference on the tendon), on the same side as the stimulus, to record the electromyographic potential (EMG). A ground electrode was placed on the sternum. The EMG was amplified and bandpass filtered (8 Hz–1.6 kHz). A 10-ohm, 15 pin FA Bone Vibrator transducer (Cadwell Laboratories, Inc., Kennewick, WA, USA, with impedance matching adaptor box) was held against the mastoid process while the child flexed his or her head against gravity to activate the SCM while looking at a picture book or playing with a toy. Preliminary trials showed that the children were reluctant to use the earphone/AC stimulation (maybe it was associated with hearing tests) though they were willing to use the bone conducted stimulus. One hundred μs clicks were generated by the Cadwell Wedge (Kennewick, WA) at 95 dB, 16 Hz. 95 dB is commonly used in the literature for air- and bone-conducted VEMP though there have been reports of a less intense stimulus (e.g., 50 dB) [14]. A consistent stimulus that was well tolerated by all participants was used in this case. The baseline EMG activity of the SCM was visualized and then 200 tracings of the evoked potential were averaged. The left side was always tested first and the procedure was repeated on the right side. The latency and amplitude of the potential was visualized and quantified using Cadwell Sierra XP software (Kennewick, WA). Based on normative data (using air-conducted VEMP) on young children [15], a normal p1 latency was in the range of 8.3–14.4 seconds and the VEMP at least 20 mV amplitude. Todd et al. [16] have shown a slightly wider range of VEMP latencies (7.5–13.9) for bone-conducted stimulation than air-conducted stimulation in adults.

Each child was evaluated using the Movement Assessment Battery for Children (Movement ABC) (Psychological Corp., London, England, 1992) to assess their proficiency in areas of manual dexterity, ball skills, and static and dynamic balances (Table 1). If the child wore cochlear implants, these were allowed during the motor testing (but not during VEMP testing). This standardized test is designed for use with children aged four to twelve years. The raw scores are converted into scaled scores in order to ascertain where the child’s performance lies in relation to the standardized sample. The sum of the three subtests provides a total impairment score. Higher scores indicate greater impairment. The test has excellent reliability (r = 0.96) and validity in normal children and children with disabilities and from populations in Europe and North America [17–19].

Student’s t-tests were used to compare the groups of subjects with a limit of P < .05 indicating significance.

<table>
<thead>
<tr>
<th>Skill categories</th>
<th>Tasks</th>
</tr>
</thead>
<tbody>
<tr>
<td>Manual dexterity</td>
<td>Insert coins in a box (with preferred and nonpreferred hands)</td>
</tr>
<tr>
<td></td>
<td>Threading beads</td>
</tr>
<tr>
<td></td>
<td>Trace bicycle trail with pen</td>
</tr>
<tr>
<td>Ball skills</td>
<td>Catch bean bag</td>
</tr>
<tr>
<td></td>
<td>Roll ball into goal</td>
</tr>
<tr>
<td>Static and dynamic balances</td>
<td>One-leg balance (preferred and non preferred legs)</td>
</tr>
<tr>
<td></td>
<td>Jump over a cord at knee level</td>
</tr>
<tr>
<td></td>
<td>Walking with heels raised</td>
</tr>
</tbody>
</table>

Table 1: Skills quantified by the Movement ABC.
3. Results

A VEMP was identified on both sides in only 4 of the HI children (3 of these children were identified as HI at birth). Four children with normal hearing of the same age as the four HI children with normal VEMP were recruited for comparison. There were no significant differences between these two groups in the total or subscores on the Movement ABC (Figure 1) though the HI children tended to have greater impairment on the manual subtest \((P < 0.07)\) and ball subtest \((P < 0.2)\) using a paired \(t\)-test.

All of the figures are scaled to the same number to allow easy comparison. It should be noted that power analysis indicated that to reject the null hypothesis correctly at the .05 level, a sample size of 14 was needed. So caution should be taken in the interpretation of this limited number of subjects though the scores indicate that all of the children in Figure 1 are functioning similarly to the standardized group.

Seven children demonstrated a VEMP on only one side (Figure 2) as measured by the action potential amplitude >20 \(\mu v\) and latency \((\text{mean} = 10.3 \pm 1.5 \text{ ms})\). There was a tendency \((P < 0.9\)–.26) for HI children with one or both sides intact \((n = 11)\) to perform better on the Movement ABC than HI children with no VEMP \((n = 23)\) (Figure 3). Of the HI children with an asymmetrical response, one child had a right cochlear implant and VEMP only on the left side, while the rest of the children had bilateral cochlear implants \((n = 2)\) or no implants \((n = 4)\).

Twenty-four children were HI since birth or within 6 months (according to the parents); nine children developed HI after the age of two years (7 at about 24 months, 2 at about 36 months). Two years was used as the natural dividing point between groups. The children with HI since birth had significantly higher scores (more developmental delay) \((P < 0.01)\) on the manual and balance subtests and the total Movement ABC (Figure 4). Two of the children in each group retained their saccular receptors bilaterally as shown by VEMP.

No significant difference was seen in the Movement ABC scores between the HI children with and without cochlear
implants (mean of the total ABC scores = 12.6 ± 9.6 and 15.5 ± 9.9, resp.).

4. Discussion

The results of this study are important to show that many but not all hearing impaired children have saccular deficit (s). A more strategic finding is the importance of the timing of the onset of the hearing impairment and probably saccular loss to the development of balance strategies.

The early postnatal period is critical for the vestibular system in experimental models such as the ferret, so that it impacts neuromuscular development of postural muscles and the development of balance strategies [8]. Human studies have focused more on motor development in the deaf population as a uniform population [12] or the effect of cochlear implants. There have been several studies of motor development of HI children with mixed results [20–24]. Some studies [20, 21] that have included HI children younger than 7 years old also include children up to 17 years of age. Rine et al. [12] noted that HI children did have delayed motor development which seemed to decelerate annually. They related this delay to “vestibular hypofunction” and supported the need for vestibular testing and repeated motor development testing during preschool and elementary school. It is unknown wether all of the vestibular receptors are lost at the same time as the cochlear receptors but the parents in this study report that the hearing loss was identified in many of the children neonatally, implying the need for very early identification of children with vestibular deficits. The current study found a significant difference in the motor development between the children with a profound hearing loss at birth and those that reportedly developed HI later with those with presumably congenital onset hearing loss demonstrating poorer performance. It is possible that vestibular receptors are more important during the initial six months to a year when the infant is learning to sit and stand. Normally children are walking by two years of age, albeit with a wide based gait. It is likely that HI children, who also have a vestibular deficit (at least a loss of saccular function), learn to depend on their vision and joint and skin sensation but it is hypothesized that the developmental milestones such as walking would be delayed without the vestibulospinal reflexes.

Nineteen of the HI children in this sample also had cochlear implants. Some of the parents reported that there were 1-2 days post implantation when the children seemed to be a little more “clumsy,” after which the children quickly returned to their presurgical motor coordination. This brief change was not measured, and at the time of this study, the children with and without cochlear implants were not significantly different in their performance of the Movement ABC. The VEMP was normal on at least one side in six (of 19) of the HI children with cochlear implants. This would seem to indicate that the cochlear implant does not have a long-term effect on balance and motor control. Cushing et al. [25] studied 40 children with profound SNHL and unilateral cochlear implants. The children ranged in age from 3 to 19 years. Saccular function (tested by air-conducted VEMP) was found in 40% of their subjects but was not correlated with balance ability when tested by the balance subset of the Bruininks-Oseretsky Test of Motor Proficiency II. In contrast, horizontal canal function correlated well with the balance score which implies that angular motion and vestibulo-ocular interaction are very important to balance. The saccule detects more linear motion so the correlation of saccule hypofunction with both delayed fine motor skills and balance skills leads one to conclude that the saccule may impact both the vestibulospinal and vestibulo-ocular reflexes though its differential function remains unclear.

A limitation of the convenience sample used in this study is that parents who may have been concerned about balance dysfunction may have volunteered their children for this study. All of these parents were eager to maximize the potential development of their children.

5. Conclusion

The onset of hearing impairment (congenital versus delayed, potentially progressive) has an impact on balance and manual skills and overall motor development. How this difference relates to the development or impairment of the vestibular receptors (in the case of the current study—saccular function) is unclear and needs further study. What is happening at the level of the cochlea may parallel what is happening at the level of the vestibular end organs. The absence of VEMP bilaterally may predict poor performance on balance and manual dexterity-related tasks although a larger sample would be required to say that this is more than a trend. The VEMP has only recently been used for screening of babies [26–28]. Further research is necessary to correlate the timing of the loss of saccular receptors with motor development in children. Early intervention could potentially optimize postural control in children who are identified with vestibular hypofunction [29].

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References


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