

COMPONENTS OF HEARING SCREENING

NOTES

There are three components of hearing screening: 1) otoscopic inspection, 2) pure tones (PTs) or otoacoustic emissions (OAEs) and 3) immittance screening. All children should be screened with all three components. (Exception: children with audiometrically documented permanent hearing loss do not need to be screened with PTs or OAEs). A child failing any portion of the screening in either ear should be rescreened (with all three components and in both ears) within 4-6 weeks. A child who fails a rescreen should be referred for further evaluation by an audiologist or a physician. (The referral process will be discussed in detail later).

The Joint Committee on Infant Hearing (JCIH) recommends the following indications for use with neonates or infants (29 days through 2 years). These indicators place an infant at risk for progressive or delayed-onset sensorineural hearing loss and/or conductive hearing loss. Any infant with these risk factors for progressive or delayed onset hearing loss who has passed the birth screen should receive audiologic screening every 6 months until age 3 years. These indicators are as follows:

- a) Parental or caregiver concern regarding hearing, speech, language, and/or developmental delay.
- b) Family history of permanent childhood hearing loss.
- c) Stigmata or other findings associated with a syndrome known to include a sensorineural or conductive hearing loss or Eustacian tube dysfunction.
- d) Postnatal infections associated with sensorineural hearing loss including bacterial meningitis.
- e) In utero infections such as cytomegalovirus, herpes, rubella, syphilis, and toxoplasmosis.
- f) Neonatal indications – specifically hyperbilirubinemia at a serum level requiring exchange transfusion, persistent pulmonary hypertension of the newborn associated with mechanical ventilation, and conditions requiring the use of extracorporeal membrane oxygenation (ECMO).
- g) Syndromes associated with progressive hearing loss such as neurofibromatosis, osteopetrosis, and Usher’s syndrome.
- h) Neurodegenerative disorders, such as Hunter syndrome, or sensory motor neuropathies, such as Friedreich’s ataxia and Charcot-Marie-Tooth syndrome.

What you hope to see during the otoscopic inspection is the eardrum. It is usually a pearly-gray color. If you see patches of white on the eardrum, you are probably seeing some scarring. This may alert you that a child has had some ear problems but it is not a cause for alarm or physician referral. Proceed with the immittance screening. An eardrum may be red or even orange-red. Again, you can note these observations on the roster but continue with the immittance screening.

An otoscopic inspection may allow you to see ventilation tubes if they are present. Pressure equalization tubes come in different colors. Some of the more common colors are silver, white, and bright green. The silver and white tubes are generally a bobbin shape, and the green ones are commonly straw-like in shape. Your immittance measurements will give you valuable information as to the status of the P.E. tube(s). Note the presence of tubes on your roster. This, in conjunction with immittance results, is useful information to the physician.

You may be able to see a perforation of the eardrum if the hole is fairly large – small perforations can be difficult to see. Immittance measurements will provide objective data as to the status of the eardrum.

Pure Tones (PT) and Otoacoustic Emissions (OAE)

Pure tone (PT) screening uses pure tones (single frequency sounds without accompanying overtones or other sounds) generated by an audiometer. PT screening has long been the technique used to screen hearing but because it requires a response from the child (e.g., hand raising; block in a bucket) it precludes the successful screening of children who are developmentally less than about three years of age.

Otoacoustic emissions (OAEs) are sounds generated within the cochlea that can be measured in the ear canal. The origin of this emission is thought to be the outer hair cells. Because it requires no response from the child it allows us to gain ear specific information about children who are developmentally unable to participate in traditional PT screening.

Both PTs and OAEs give us information about each ear but because pure tones can be presented at lower levels than OAE screening equipment uses, they should be used as soon as the child is developmentally capable of participating. There is no

magical age at which this occurs but a guideline is a developmental age of three years. At that age attempt PTs and if unsuccessful, use OAEs.

NOTES

Hearing screening in sound field (using loud speakers) is not appropriate for three reasons. First, it does not give ear specific information; it tests only the better ear if there is a difference between ears (like taking an eye test with both eyes).

Considering the evidence we have about UHL it is essential that we know the status of both ears. Secondly, while using a loud speaker, it is difficult to determine if a child is responding to an auditory stimuli or simply moving around. In an environment other than a sound-treated room it is very easy to misinterpret a child's responses. Lastly, calibration of sound field equipment is problematic.

Otoacoustic Emissions (OAE) Screening

When doing OAE screening, choose a quiet room. Infants are best screened when they are in a quiet state. Visual stimuli such as a movie, quiet toys, books, computer games, etc. may be useful in diverting the child's attention from the screening. Also, giving the child something to hold in each hand may be useful in keeping wandering hands from removing the probe from their ear. An otoscopic inspection will help you determine an appropriate sized probe tip. Once the probe is in place and the child is quiet and calm, the screening may begin. **The most frequent cause of OAE failure is a poorly fit probe.** Quality of the probe fit is the first and most important factor in collecting accurate data. It is often difficult to achieve a perfect fit on the first attempt; refitting commonly is necessary. The fit of the probe in the canal is crucial to the presentation of an optimal stimulus and also to sealing off ambient noise that may contaminate the measurement.

The most common OAE screening devices currently used in Wyoming Child Development Regions are the Biological AuDX and the ECHOCHECK. Each will be discussed in terms of their operation and failure criteria.

AuDX Procedures

- 1) Connect the probe assembly to the AuDx box.
- 2) Place the other end of the probe assembly with the appropriate sized probe tip into the child's ear canal.
- 3) Press the ON button.
- 4) Press the DX (SELECT) button to continue.
- 5) When PERFORM DPOAE appears in the display, press

Check the earphone cushions; they should be reasonably soft, resilient, and free from cracks. Occasionally, they should be wiped clean with warm water and mild soap. Check the shape of the headband; the earphone cushions should lie together with a small amount of tension. If the cushions do not meet or if they seem to lie together with too much tension, shape the headband by bending it with a twisting motion.

Any extraneous noise present in the earphones may affect screening results. To check for this, set the frequency at 1000 Hz and the intensity level at 90 dB. Interrupt the tone and listen for noise. Listen again with the instrument at 60 dB and again at the lowest setting of the attenuator (hearing level dial). If humming or extraneous noises are heard at these settings, the instrument should not be used until the problem is addressed.

The earphone cords should be checked for breaks and loose connections. Set the attenuator at 40 dB and 2000 Hz. While listening to the tone, flex the cord along its length and especially at its connections (at earphones and audiometer). If a scratchy noise is heard or the tone is intermittent discontinue screening until the cords have been replaced or the connections are made secure. Under no circumstances should headphones from one audiometer be interchanged for headphones of another audiometer. Earphones are calibrated as an integral part of the instrument. Earphones cannot be interchanged even temporarily without conducting calibration using a sound level meter.

Lack of lubrication or the presence of dirt deposits in the attenuator may cause noise to be heard in the earphones when the hearing levels are changed. In order to check for this, set the frequency to 250 Hz and then slowly increase the attenuator from 0-60 dB and listen for scratchy noise. If the noise is detected, the instrument may be used, if necessary, as long as adjustments of the attenuator are made only when the tone is off.

Scratchy noises or clicks may develop in the frequency switch. To check this, set the attenuator at 30 dB and slowly increase the frequency from low to high. If noise is present, you can prevent the noise from getting to the headphones by changing the frequency dial only when the tone is off. With the newer digital equipment, noises and clicks from the attenuator and frequency dials is not a common problem.

All of the preceding checks are made with the earphones properly placed on the adult screener and with the assumption that s/he has normal hearing for 250 Hz through 8000 Hz.

performs the measurements automatically once an adequate seal is obtained by holding the probe tip at the entrance of the external ear canal. The manipulation of the TM-middle ear system is accomplished pneumatically (i.e., via air) and is known as tympanometry.

The acoustic reflex (AR), also known as the stapedial reflex, is the contraction of one or more muscles of the middle ear (stapedius and tensor tympani) in response to the presentation of a loud stimulus. This contraction causes a stiffening of the ossicular chain which in turn “stiffens” the tympanic membrane. An AR is not observable in middle ears with effusion or other tympanic membrane and/or middle ear conditions causing stiffness.

Many immittance screening bridges have the AR feature and they provide a graphic representation of the muscle contraction. This is interpreted as a “present AR”.

As a screening tool the presence of the AR can help clarify questionable tympanograms as in the case of low (.1) compliance but with an observable peak. No interpretation can be made when a screening AR is absent because an absent AR may be caused by too little stimulation, instrumentation artifact, or its absence may be normal for a particular ear. However, the presence of an AR rules out significant middle ear pathology.

Because the middle ear system is mechanical, immittance measurements can be made objectively. Children do not need to respond in any way and the test can be performed in a noisy environment.

Tympanometry should not be performed on infants less than 7 months of age. The ear canals of infants in this age group are very pliable and immittance measurements may be the result of pneumatically moving the ear canal wall rather than the TM.

Tympanometry is conducted automatically when the immittance equipment (also called an immittance bridge) creates in sequence, positive and negative air pressures in a sealed ear canal and records the changes caused in the mobility of the tympanic membrane/ossicular chain system. This changing relationship is plotted as a tympanogram.

Positive air pressure created in the ear canal increases the stiffness of the TM-middle ear system and reduces its compliance (mobility). This is the starting point of the tympanogram (“A,” Fig. 4). As pressure is progressively lowered, compliance is altered. When air pressure on both sides of the TM is equal, a peak on the tympanogram tracing will

Ear canal volume, also called physical volume (PV) or equivalent canal volume is obtained when an air pressure of +200 daPa is applied to the ear canal and “clamps” the TM. PV is simply a measurement of the amount of space between the probe tip and “wherever the air can’t get any farther.” Volume measurements are helpful in determining if ventilation tubes are patent (open), if the TM has a perforation, or if the ear canal is occluded with wax or other foreign objects.

Equipment

Immittance screening equipment provides a visual representation of the tympanogram and most display numeric values for MEP, TM compliance, PV, and TW. Some instruments measure TM compliance and PV in cubic centimeters (cc) and some use milliliters (ml). This manual will use ml, but practically speaking, they are interchangeable. Older equipment measures MEP in mmH₂O but current models use decaPascals (daPa). This manual uses daPa but, again, the terms are essentially interchangeable.

Immittance screening equipment is designed to be completely automatic; i.e., once it starts it doesn’t stop, no matter what! The result is a measurement for every aspect (MEP, PV, TM compliance, and TW). **By knowing the significance of each measurement and their relationship to each other** you can determine the validity of the values and make appropriate decisions from your screening results.

A daily check of your immittance bridge is recommended. First, calibrate the unit according to the manufacturer’s specifications. Next, run a tympanogram on yourself. One of the most common problems with an immittance bridge is a blocked probe assembly. Consult the manual about cleaning the probe and follow the directions carefully and regularly. Probe assemblies can be replaced but they are expensive.

Criteria

The failure criteria for immittance screening for children is

- o MEP > -250 daPa
- o TM compliance < .2 ml. (Note: If the compliance is .1 AND a peak is observed AND an acoustic reflex is present it is considered a PASS.)

The estimated equivalent ear canal volume measures of preschool children prior to and following placement of tympanostomy tubes (ECV for most ears with and without tubes)

- o Pretube 0.3 – 0.9 cm³
- o Post-tube 1.2 – 5.5 cm³

	1 st grade	2 nd grade	3 rd grade	5 th grade
n	80	82	90	80
Range	1.0 - 2.1	1.0 - 1.9	1.0 - 2.3	1.1 - 2.2
Range (minus lowest and highest value)	1.1 - 1.9	1.2 - 1.8	1.1 - 2.2	1.2 - 2.1
Mean	1.46	1.501	1.592	1.667
Mode	1.5	1.5	1.5	1.6; 1.9

Ear canal volumes (in ml) for students in 1st, 2nd, 3rd, and 5th grades.

Previously, abnormal physical volume (PV) was used as a failure criteria for immittance screening but because of the huge variability among children it is difficult to determine what is “too large” or “too small.” PV must be viewed in relationship to the compliance and MEP values and by doing so you can estimate the status of the middle ear (ME) system. (Is the eardrum intact? Is the tube patent? Is there a perforation? Is the ME system normal?)

The use of PV, in isolation, as failure criteria is problematic because of the huge variability among children. What is important to know (and what will be discussed in the next chapter) is the relationship of PV to other aspects of immittance screening. This knowledge will assist you in making appropriate referrals from your hearing screening program.

Any child failing TM compliance or MEP measurements should be rescreened (with all three components of hearing screening in both ears) within 4-6 weeks. Children failing the immittance rescreen should be referred for medical evaluation.

Procedure

To conduct immittance screening, position the child in such a way that you can visually inspect the external ear canal for size and direction of the ear canal. With preschoolers try sitting in an adult-size chair while having the child stand in front of you. For infants and toddlers have a familiar adult hold the child on their lap. Select an appropriate sized probe tip for the size of the ear canal opening. The probe tip should be larger than the opening of the external ear canal. The intent is to obtain an airtight seal at the canal opening; the probe tip should NOT be inserted into the canal itself. Fit the base of the rubber probe tip over the end of the metal or plastic probe stem assembly on the immittance bridge so that it goes completely over the probe stem. Generally, the same size tip is appropriate for a similar age group. Clean the probe tip with an alcohol wipe after each child.

Getting and maintaining a good seal at the opening of the external ear canal opening is the most important variable to accurate and reliable immittance screening. In order to obtain the best external ear canal seal, gently pull the top of the test ear up and back to straighten the ear canal. Apply the probe tip firmly to the canal opening. Depending on the unit you are using, you may see a flashing light or a steady green light as long as the seal is being maintained. Some immittance bridges will display 'air leak', 'block', or 'testing' on the display screen depending upon the status of the external ear canal seal. If you are having difficulty maintaining a seal, it is best to remove the probe, reexamine the direction of the canal, and try again. Changing the size of the probe tip may help maintain an external ear canal seal. If the child fails the immittance screening, immediately repeat the measurement.

The directions you give to the child should be very simple; all they really need to do is stand (sit) quietly. Sometimes they will ask, "What's that?" Assuming you have already done an otoscopic inspection, you can respond with, "It looks at your ear like I did." Occasionally, a child might ask "What does it do?" "It checks your eardrum," is an appropriate response. Don't forget to praise and encourage the young child while doing immittance measurements. If the child is 'squirmy' you will be able to complete the measurements by talking to him/her as the test is performed. If your immittance unit has lights, you can simply ask the child to watch the lights. Some children also enjoy watching the tracing being completed.

Screeners have reported a variety of techniques and situations, which help divert an infant, toddler or preschooler's attention away from the screening procedures thus enabling the screening process to be completed. Some of their "tricks" include having the child watch themselves in a mirror, providing a TV/VCR cartoon for the child to watch, or allowing the child to scribble on a piece of paper.

Immittance Measurements As They Relate To Disorders

In the case of atresia, you will probably not be able to obtain immittance measurements simply because there usually is no canal opening. If there is an opening, your immittance results will probably be low compliance (there may be no eardrum to pneumatically move) and possibly a small external ear canal volume measurement (depending on where the malformation of the ear canal begins).

You may not be able to visualize the eardrum because of wax, but if the air from the immittance bridge can get around the wax, you can still get a tympanogram. If your otoscopic inspection revealed excessive cerumen, it still is appropriate to perform the immittance screening. If your otoscopic inspection revealed a foreign object (this does not include excessive wax), stop the screening and refer the child to their physician for removal of the foreign object.

The presence of a poorly functioning Eustachian tube is indicated by negative middle ear pressure (MEP). The significance of negative MEP is that it occurs before effusion develops and again as effusion resolves. Normally, MEP is at or near 0 daPa. When the Eustachian tube is not functioning adequately negative middle ear pressure occurs. If this negative middle ear pressure persists for a period of time, the vacuum which is created in the middle ear cavity may begin to draw moisture from the mucosal lining in the middle ear cavity and fluid (effusion) may form. As the effusion resolves, negative middle ear pressure is again seen before the middle ear pressure returns to, or near, 0 daPa.

Tympanic membrane compliance may be normal in the presence of negative MEP but as the MEP becomes more negative the compliance decreases and the TW increases. In other words as the TM is 'pulled in' by the negative MEP it becomes less mobile and the shape of the tympanogram becomes shorter and wider. Low compliance produces a screening failure and thus identifies the child with possible middle ear problems.

Check the manual for your immittance bridge to find out the pressure range of your equipment. Most are from +200 to approximately -300 daPa. If the child's MEP is in excess of that range the unit may display a question mark or other indicator rather than a numerical value. This doesn't mean the child doesn't have a MEP, it simply means the equipment can't find it.

Although unusual, you may see positive MEP. A reading in excess of +100 daPa can also reflect an error in the immittance procedure. If this occurs immediately repeat the measurement.

When fluid is present in the middle ear space the tympanogram will be flat. The tracing may 'bounce' along the bottom of the scale or you may see a rounded tracing without a peak. The immittance bridge will display a compliance value $< .2$ ml. Most equipment will be unable to specify a MEP value (the unit cannot identify a point of maximum compliance) and MEP will be identified as '?' or No Peak (NP). This flat tracing occurs because the fluid behind the TM produces resistance or 'pushes' from that side of the TM, which prevents air (via tympanometry) from moving the eardrum from the other side.

Again, it is the relationship of the PV to the other immittance values that determines its significance. PV is irrelevant in the presence of normal TM compliance and normal MEP but significant when MEP and TM compliance are abnormal. It is critical for you to understand how your particular piece of equipment (immittance bridge) obtains and displays data (results).

When it is necessary to communicate screening results to parents or physicians either verbally or in writing keep in mind that immittance screening does not allow a medical diagnosis to be made – that is done by the physician. Appropriate phrases include: “Immittance measurements suggest....”; “Results are consistent with...”; “...support the presence of...”

THE AUDIOGRAM

This section is included to help you interpret audiograms and reports you receive from other facilities. A basic knowledge of the audiogram should facilitate your communication with audiologists and with families and teachers.

An audiogram is a graphic representation of a person’s hearing at a specific point in time. For children with conductive hearing loss, audiograms can be different from one evaluation to the next due to fluctuations in hearing. As you use the IEP/IFSP process to plan for children you should have current audiological information. For infants with sensorineural hearing loss hearing should be evaluated every 3-4 months for the first year after identification, at least twice a year through the preschool years and, at a minimum, annually after entering school. The purpose of close audiological monitoring is to identify progressive hearing loss.

In the past young children were tested in sound field. The child sat on a parent’s lap inside a sound proof booth with a speaker on either side. As stimuli were presented via the speakers and the child turned toward the signal she/he was reinforced with a blinking light. The technique, called Visual Reinforcement Audiometry (VRA) does not provide ear specific information; it only tests the better ear if in fact there is a difference. If you are referring a child because they failed the OAE component of the hearing screening **you should expect and request ear specific information from the audiological assessment.**