### Audiological Procedures to Be Utilized During Data Collection for Objective 3

<table>
<thead>
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<th>Audiological Test, Procedure and Duration</th>
<th>Motivation</th>
<th>Tool</th>
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<tr>
<td>Interview</td>
<td>The purpose of obtaining case history information will be to aid in establishing an overview of the participant’s auditory status, medical conditions, medication, noise exposure and communicative abilities related to audition [1]. Information on the risk factors for ototoxic hearing loss will, thus, also be documented during the case history interview. This questionnaire will also include questions on tinnitus, as a common complaint of cancer patients on chemotherapy is the onset of tinnitus.</td>
<td>Case history form, focusing on hearing history, medical history, family history, and history of noise exposure, was developed. Questions on tinnitus were adapted from the Tinnitus Otoxicity Monitoring Interview (TOMI) schedule [2].</td>
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<td>Otoscopic Examinations</td>
<td>An otoscopic examination of both ears will be conducted to determine the status of the tympanic membrane and the external ear [3].</td>
<td>An Agine otoscope will be used to conduct otoscopic examinations.</td>
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<td>Imittance Audiometry-Tympanometry and ipsilateral and contralateral acoustic reflex threshold testing</td>
<td>Tympanometry will be conducted to determine if there are any abnormalities in the middle ear as well as to aid in detection and differentiation of possible pathologies [1]. The acoustic reflex threshold is the lowest intensity of a sound stimulus that elicits a measurable change in the acoustic admittance [4]. “These measures provide information related to the function of the middle ear and the sensory, neural, and motor pathways associated with the reflex arc” [5] (p. 169). Both ipsilateral and contralateral tympanometry will be conducted.</td>
<td>A clinical impedance audiometer, the GSI Tymstar V2 Impedance meter will be used for immittance audiometry.</td>
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be placed in the entrance of the participant’s ear canal and will introduce slight pressure as well as soft beeping sounds. This test does not cause any discomfort and the participant will not be required to respond. **Duration:** ± 5 minutes

| **Pure Tone Audiometry** (air and bone conduction) |  
| **Procedure:**  
Headphones will be placed on the participants’ ears and they will hear a beeping sound, where they will be required to press a button to indicate that they heard the sound. |  
| **Duration:** ± 20 minutes |  

Since pure tone audiometry is the foundation of every audiological evaluation, its’ results will form the basis from which an initial diagnosis of hearing loss can be made [6]. It will be conducted to also quantify the degree of hearing loss and determine the type of hearing loss [7].

Pure tone air conduction thresholds will be obtained bilaterally via air conduction at the following frequencies: \(125, 250, 500, 1000, 2000, 4000, 6000, 8000, 9000, 10000, 11200, 12500, 14000, 16000, 18000\) and \(20000\) Hz, while bone conduction thresholds will be obtained bilaterally at \(250, 500, 1000, 2000\) and \(4000\)Hz.

Bone conduction audiometry will be conducted at baseline, post treatment audiometry 6 months after completion of chemotherapy and if there is a significant change in the air conduction thresholds during the other audiological evaluations.

A twin channel clinical diagnostic audiometer, the Madsen Astera will be used for pure tone audiometry.
**Speech Audiometry**

**Procedure:** Headphones will be placed on the participants’ ears and they will hear words which they will be required to repeat.

**Duration:** ± 10 minutes

Speech recognition threshold (SRT) testing will be conducted to determine the lowest hearing level at which spondaic words are identified 50% of the time \[3\]. It is also a tool used for the confirmation of pure tone thresholds and in doing so, alerts the audiologist to invalid pure tone results \[8\].

Speech recognition score testing is conducted to measure how well the listener can understand speech as a function of the ability to differentiate sounds under optimum circumstances. The score is intended to be a measure of the clarity with which the patient hears speech \[8\]. Speech audiometry will be conducted for the baseline audiological evaluation and the 6 month follow-up evaluation.

**Distortion Product Oto-Acoustic Emission Testing**

**Procedure:** A soft probe will be placed in the entrance of the participant’s ear canal and will introduce soft beeping sounds. This test does not cause any discomfort and the participant will not be required to respond.

**Duration:** ± 5 minutes

“OAEs are believed to be the by-products of the preneural mechanisms of the cochlear amplifier, and in particular to be linked to the normal functioning of the outer hair cells”\[10\] (p. 441). Therefore, OAEs are sensitive to hearing losses, resulting from outer hair cell damage. OAE results will help with the differentiation of cochlear vs. retrocochlear disorders and also identify individuals with subtle abnormalities of CNS function \[11\].

SRT- The CID Spondee word list will be used. For isiZulu speakers, the digits test, will be used, as it is low linguistically loaded \[9\].

SRS- The CID W-22 Auditory test word list will be used. For isiZulu speakers, an isiZulu wordlist collated in the Discipline of Audiology, will be used. The audiometric sound proof booth and the twin channel audiometer, as described for pure tone audiometry will be used.

The Mico Oto–acoustic emissions will be used for the elicitation of the OAE.
References


