

**Ball State University**  
**Audiology Clinic**  
**Adult Case History**

Name \_\_\_\_\_ Birth Date \_\_\_\_\_ Age \_\_\_\_\_ Gender  M  F

Audiologist \_\_\_\_\_ Grad. Clinician \_\_\_\_\_ Date \_\_\_\_\_

Referral Source \_\_\_\_\_

Occupation \_\_\_\_\_

1. **Reason for Visit:** \_\_\_\_\_

\_\_\_\_\_

2. **Hearing Loss:**  Yes  No Approx year or age of onset: \_\_\_\_\_

Which ear:  RT  LT  Both Better ear:  RT  LT  Unknown

Hearing loss characteristics:  Rapid onset/changes  Gradual onset/changes  Fluctuating

Family history:  Yes  No Who: \_\_\_\_\_

Situations that cause difficulty: \_\_\_\_\_

Remarks: \_\_\_\_\_

3. **Previous Hearing Evaluation:**  Yes  No

When / Where: \_\_\_\_\_

Results: \_\_\_\_\_

Recommendations made: \_\_\_\_\_

4. **Noise Exposure:**  Yes  No

Types of noise:  Occupational: \_\_\_\_\_

Recreational: \_\_\_\_\_

Other: \_\_\_\_\_

Hearing protection used?  Yes  No Percentage of time: \_\_\_\_\_

Remarks: \_\_\_\_\_

5. **History of Ear Infections:**  Yes  No  RT  LT  Both

Frequency: \_\_\_\_\_

Treatment:  Antibiotics  PE Tubes (# of sets: \_\_\_\_\_)  Other: \_\_\_\_\_

6. **Ear Surgery:**  Yes  No  RT  LT

Date / type of surgery: \_\_\_\_\_

7. **Tinnitus:**  Yes  No      Approx year or age of onset: \_\_\_\_\_  
 RT  LT  AU / Non-localized  Constant  Fluctuating

Description: \_\_\_\_\_

Concurrent symptoms:  Dizziness  Hearing loss  Other: \_\_\_\_\_

8. **Dizziness:**  Yes  No       Vertigo  Light headed

Describe: \_\_\_\_\_

Previously evaluated?  Yes  No      Where/how: \_\_\_\_\_

Successfully treated  Yes  No      Where/how: \_\_\_\_\_  
(If patient reports dizziness, refer to the Dizziness / Balance Case History)

9. **Other Auditory Symptoms:**

Aural fullness/pressure ( RT  LT)  Hypersensitivity to loud sounds

Otagia ( RT  LT) Describe: \_\_\_\_\_

Other: \_\_\_\_\_

10. **Hearing Instruments:**  Currently uses  Previously used  Recommended, never used

Ear fit:  RT  LT  AU      Style: \_\_\_\_\_

Perceived benefit: \_\_\_\_\_

Comments: \_\_\_\_\_

11. **General Health:**

Good  Other: \_\_\_\_\_

Current medications: \_\_\_\_\_

\_\_\_\_\_  
\_\_\_\_\_

12. **Comments / Additional Information:**

\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

# Ball State University Speech and Audiology Clinic – Family Medical History Form

Patient: \_\_\_\_\_ Date: \_\_\_\_\_ Birthdate: \_\_\_\_\_ Sex: \_\_\_\_\_

Address: \_\_\_\_\_ City: \_\_\_\_\_ State: \_\_\_\_\_ ZIP: \_\_\_\_\_

Home Phone: ( ) \_\_\_\_\_ Other Phone: ( ) \_\_\_\_\_ Email: \_\_\_\_\_

Primary Care Physician: \_\_\_\_\_ City: \_\_\_\_\_

Maternal Ethnicity of Patient: \_\_\_\_\_ Paternal Ethnicity of Patient: \_\_\_\_\_

Person Completing this Form (if not patient): \_\_\_\_\_ Relation to patient: \_\_\_\_\_

Reason for Appointment: \_\_\_\_\_

Please indicate if the patient or any relatives (including at least 3 generations) currently experience or have experienced the listed symptoms or conditions. Please provide additional information if possible and note which relative (e.g., maternal grandmother) when applicable. Information on family health can be useful in understanding potential causes for hearing, speech or language difficulties or other health issues. While some symptoms (e.g., miscarriages or fainting spells) may seem irrelevant, certain combinations of symptoms may signal an underlying condition that is related to the patient’s primary concern.

**Otologic and Hearing problems:**

**Relationship/Comments:**

- | Patient                  | Relative                 |  |
|--------------------------|--------------------------|--|
| <input type="checkbox"/> | <input type="checkbox"/> | Hearing Loss Present at Birth, Childhood, or Young Adulthood _____ |
| <input type="checkbox"/> | <input type="checkbox"/> | Hearing Loss, Any Age of Onset _____                               |
| <input type="checkbox"/> | <input type="checkbox"/> | Vertigo/Dizziness/Balance problems _____                           |
| <input type="checkbox"/> | <input type="checkbox"/> | Other _____  |

**Vision Problems:**

- | Patient                  | Relative                 |   |
|--------------------------|--------------------------|---|
| <input type="checkbox"/> | <input type="checkbox"/> | Premature and/or Progressive, Significant Vision Loss _____ |
| <input type="checkbox"/> | <input type="checkbox"/> | Night Blindness _____                                       |
| <input type="checkbox"/> | <input type="checkbox"/> | Tunnel Vision _____   |

**Learning Disabilities/Developmental Delays:**

- | Patient                  | Relative                 |  |
|--------------------------|--------------------------|--|
| <input type="checkbox"/> | <input type="checkbox"/> | Cognitive Function Impairment (Mental Retardation) _____ |
| <input type="checkbox"/> | <input type="checkbox"/> | Speech/Language Delay _____                              |
| <input type="checkbox"/> | <input type="checkbox"/> | Motor Delay _____  |
| <input type="checkbox"/> | <input type="checkbox"/> | Learning Disability of Unknown Cause _____               |
| <input type="checkbox"/> | <input type="checkbox"/> | Autistic Spectrum Disorders _____                        |
| <input type="checkbox"/> | <input type="checkbox"/> | Other _____  |

**Neurological or Cardiovascular Concerns:**

- | Patient                  | Relative                 |                           |
|--------------------------|--------------------------|---------------------------|
| <input type="checkbox"/> | <input type="checkbox"/> | Seizures _____            |
| <input type="checkbox"/> | <input type="checkbox"/> | Fainting _____            |
| <input type="checkbox"/> | <input type="checkbox"/> | Tremors _____             |
| <input type="checkbox"/> | <input type="checkbox"/> | Heart Defect _____        |
| <input type="checkbox"/> | <input type="checkbox"/> | Heart Attack _____        |
| <input type="checkbox"/> | <input type="checkbox"/> | Irregular Heartbeat _____ |

- |                          |                          |                                  |
|--------------------------|--------------------------|----------------------------------|
| Patient                  | Relative                 |                                  |
| <input type="checkbox"/> | <input type="checkbox"/> | Sudden Death (unexplained) _____ |
| <input type="checkbox"/> | <input type="checkbox"/> | SIDS _____                       |

**Musculoskeletal**

- |                          |                          |  |
|--------------------------|--------------------------|--|
| <input type="checkbox"/> | <input type="checkbox"/> | Unusual Stature _____                    |
| <input type="checkbox"/> | <input type="checkbox"/> | Craniofacial or Oral Abnormalities _____ |
| <input type="checkbox"/> | <input type="checkbox"/> | Cleft Palate/Lip _____                   |
| <input type="checkbox"/> | <input type="checkbox"/> | Ear Anomalies _____                      |
| <input type="checkbox"/> | <input type="checkbox"/> | Arthritis _____                          |
| <input type="checkbox"/> | <input type="checkbox"/> | Frequent Broken Bones _____              |

**Kidney**

- |                          |                          |                           |
|--------------------------|--------------------------|---------------------------|
| <input type="checkbox"/> | <input type="checkbox"/> | Kidney Malformation _____ |
| <input type="checkbox"/> | <input type="checkbox"/> | Kidney Disease _____      |
| <input type="checkbox"/> | <input type="checkbox"/> | Blood in Urine _____      |

**Pigmentation**

- |                          |                          |  |
|--------------------------|--------------------------|--|
| <input type="checkbox"/> | <input type="checkbox"/> | Heterochromia (eyes of different color or a multi-colored eye) _____ |
| <input type="checkbox"/> | <input type="checkbox"/> | Hair (early graying, white patches) _____                            |
| <input type="checkbox"/> | <input type="checkbox"/> | Vitiligo _____   |
| <input type="checkbox"/> | <input type="checkbox"/> | Café au Lait Spots _____   |

**Reproductive/Pregnancy Concerns:**

- |                          |                          |  |
|--------------------------|--------------------------|--|
| Patient                  | Relative                 |  |
| <input type="checkbox"/> | <input type="checkbox"/> | Stillbirth _____                               |
| <input type="checkbox"/> | <input type="checkbox"/> | Miscarriage _____                              |
| <input type="checkbox"/> | <input type="checkbox"/> | Early Menopause _____                          |
| <input type="checkbox"/> | <input type="checkbox"/> | Consanguinity (interfamily relationship) _____ |
| <input type="checkbox"/> | <input type="checkbox"/> | Infertility _____                              |

**Endocrine:**

- |                          |                          |  |
|--------------------------|--------------------------|--|
| Patient                  | Relative                 |  |
| <input type="checkbox"/> | <input type="checkbox"/> | Diabetes _____                                     |
| <input type="checkbox"/> | <input type="checkbox"/> | Thyroid Disorder (enlarged thyroid - goiter) _____ |

**Genetic Disorders & Syndromes:**

- |                          |                          |  |
|--------------------------|--------------------------|--|
| Patient                  | Relative                 |  |
| <input type="checkbox"/> | <input type="checkbox"/> | Stickler Syndrome _____                      |
| <input type="checkbox"/> | <input type="checkbox"/> | Usher Syndrome _____                         |
| <input type="checkbox"/> | <input type="checkbox"/> | Branchio-oto-renal Syndrome _____            |
| <input type="checkbox"/> | <input type="checkbox"/> | Pendred Syndrome _____                       |
| <input type="checkbox"/> | <input type="checkbox"/> | Neurofibromatosis type II _____              |
| <input type="checkbox"/> | <input type="checkbox"/> | Mitochondrial Disorders _____                |
| <input type="checkbox"/> | <input type="checkbox"/> | Alport Syndrome _____                        |
| <input type="checkbox"/> | <input type="checkbox"/> | Waardenburg Syndrome _____                   |
| <input type="checkbox"/> | <input type="checkbox"/> | Long QT Syndrome/Jervell Lange-Nielsen _____ |

# Ball State University Speech and Audiology Clinic – Family Medical History Form

## Genetic Disorders & Syndromes *(continued)*:

## Relationship/Comments:

Patient	Relative	
<input type="checkbox"/>	<input type="checkbox"/>	Pierre-Robin Sequence _____
<input type="checkbox"/>	<input type="checkbox"/>	Treacher Collins Syndrome _____
<input type="checkbox"/>	<input type="checkbox"/>	Connexin 26 Deafness _____
<input type="checkbox"/>	<input type="checkbox"/>	Other Syndromes or Chromosomal Abnormalities _____

Has a family member ever had an evaluation by a geneticist?  yes  no

Please make any additional comments in the space provided below.

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