

Ball State University
Audiology Clinic
Child Case History

Name _____ Birth Date _____ Age _____ Gender M F

Audiologist _____ Grad. Clinician _____ Date _____

Referral Source _____ Child's PCP _____

School _____ Grade _____ Teacher _____

1. Reason for Visit / Parental Concern: _____

2. Previous Hearing Evaluation: Yes No

When / Where: _____

Results: _____

Recommendations made: _____

3. History of Ear Infections: Yes No RT LT Both

Last occurrence: _____ First occurrence: _____

Frequency: _____

Treatment: Antibiotics PE Tubes (# of sets: _____) Other: _____

4. Hearing Aids: Yes No

Ear fit: RT LT AU Make / Model: _____

Age when first fit: _____ Hours used per day: _____

5. Prenatal, Birth, and Developmental History

a. Risk Factors Check if no risk factors

Family History of Permanent Childhood
Hearing Loss

Mechanical Ventilation

Craniofacial Anomalies

Cytomegalovirus (CMV)

Bacterial Meningitis

In utero Infection

Rubella

Special Care / NICU (over 5 days)

Herpes

Hyperbilirubinemia requiring exchange or transfusion

Toxoplasmosis

Ototoxic Medications

Genetic Syndromes Associated with Hearing Loss

Syphilis

Extracorporeal Membrane Oxygenation (ECMO)

Other

Comments / Additional Information: _____

- b. Complications During Pregnancy None

Describe: _____

- c. Alcohol / drug use during pregnancy (prescription or non-prescription) Yes No

Describe: _____

- d. Normal delivery? Yes No

Issues: _____

- e. Complications immediately following birth or during first two weeks Yes No

Describe: _____

- f. Significant medical / health issues since birth Yes No

Describe: _____

6. Speech / Language History

a. Age of first word: _____ First word: _____

b. How does child make his/her needs known: _____

c. Is speech understood by others Yes No

7. Developmental Milestones

a. Developmental milestones met as expected Yes No

Comments: _____

b. Is child receiving therapy from other discipline (SLP, PT, OT, etc.) Yes No

Comments: _____

8. 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.
