

## PRIOR AUTHORIZATION for HEARING AIDS

For authorization, please complete this form, include patient chart notes to document information and FAX to the PEHP Prior Authorization Department at (801) 366-7449 or mail to: 560 East 200 South Salt Lake City, UT 84102. If you have prior authorization or benefit questions, please call PEHP Customer Service at (801) 366-7555 or toll free at (800) 753-7490. *Please be aware that not all employer groups have the same hearing aid coverage.												
Section I: PATIENT INFORMATION												
Name (Last, First MI):				DOB:	DOB: A			PEHP	ID #:			
Section II: PROVIDER INFORMATION												
Date Requested:			Ordering Provider/Physician:			Ordering Provider/Physician NPI #:						
Ordering Provider/Physician Contact Person:			Phone: ( )			Facsimile:						
Rendering Provider: Renderi		Rendering	g Provider NPI #: Rendering		Provider Tax ID #: Rende		Renderin	lering Provider Address:				
Rendering Provider Contact Person:			Rendering Provider Phone:				Rendering Provider Fax:					
( )     ( )       Section III: PRE-AUTHORIZATION REQUEST												
Nat	ure of Request: <i>Please check.</i>											
Nature of Request: Please check.       Requested Date of Service:         □ Auth Extension       □ Pre-Auth       □ Retrospective Auth       □ Urgent												
Medical Diagnosis/ICD-10 Code (s):         Treating Diagnosis/ICD-10								de (s)	:			
Service (s) Requested: Please list all requested services/CPT codes regardless of pre-auth requirement.         Procedure/Service:												
Procedure/Service: CPT/HCPCS code: CPT/HCPCS code:CPT/HCPCS code:												
Procedure/Service:												
A. Does the patient currently own hearing aids?		aring aids?	A. 1. Purchase Date:		A. 2. Type of Hearing Aid		ring Aid:	id: A. 3.		Hearing Aid Condition:		
B. What type of hearing aid is being requested?												
	1. Air Conduction											
	2. D Bone Conduction											
	<ol> <li>General Fully Implantable Middle Eau</li> <li>Partially Implantable Magne</li> </ol>			hono® Alpha	2™ System	Cochleau	r™ BAHA® /	1 Attr	act)			
	5. Non-implantable, intraoral b					Cocilicai			actj			
6. □ Semi-Implantable Middle Ear (e.g. Vibrant Soundbridge, Maxum <sup>™</sup> )												
	7. Other ( <i>please specify</i> ):											
(Please check service being requested.)			QUESTION				YES	NO	COMMENTS/NOTES			
	Did any of the following conditions of a. □ Congenital hearing loss	ause the pat	ient's hearing los	s? Please c	heck all that	apply.						
<ul> <li>b. D Direct physical trauma affecting the middle ear or inner ear</li> </ul>												
c.  Tumor affecting the middle or inner ear												
d.  Radiation therapy												
<ul> <li>e. □ Infection (e.g. rubella, herpes simplex) resulting in damage to the middle ear or inner ear</li> <li>f. □ Side effect of medication (e.g., aminoglycosides, chemotherapy drugs)</li> </ul>												
<ol> <li>Is the hearing loss permanent?</li> </ol>												
3. Does hearing testing reveal hearing loss of more than 10dB across at least two frequency ranges?										Please include copy of hearing testing report.		



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Name (Last, First MI):	DOB:	Age:	PEHP ID #:		
(Please check service being requested.) QUESTION (cont'd)			YES	NO	COMMENTS/NOTES
4. Is the hearing aid expected to restore equal or greater than 259 has a 10dB loss, 2.5dB or more will be restored)?					
5. Is the hearing loss caused by natural hearing loss associated wi					
6. Is the hearing loss caused by noise exposure?					
7. Does the patient have conductive hearing loss unresponsive to					
8. Does the patient have sensorineural hearing loss?					
9. Does the patient have mixed hearing loss?					
10. D Air Conduction Hearing Aid:					
a. Is "Behind the Ear" (BTE) device being requested becau hearing loss?					
b. Is "In the Ear" (ITE) or "Completely in the Canal" (CIC) d has mild to moderate hearing loss?	he patient				
c. Is "In the Ear Canal" (ITC) device being requested becau hearing loss?	re				
d. Is "On the Body" hearing aid being requested because t	nd hearing				
e. Is "Contralateral Routing of Sound" (CROS) device being side hearing loss?	has single-				
11. D Bone Conduction Hearing Aid:					
<ul> <li>Does the patient have malformation of the external or r atresia, small ear canals that precludes the use of a conv</li> </ul>					
<ul> <li>b. Does the patient have a condition involving chronic mide chronic otitis media) that precludes the use of a convent</li> </ul>					
12. D Semi-Implantable Middle Ear Hearing Aid:					
a. Is the patient 18 years or older?					
b. Does the patient have moderate to severe sensorineura					
c. Does the patient have any of the following medical conc conduction aid? <i>Please check all that apply.</i>					
Chronic otitis externa that is unresolved despite multiple	tiple rounds of drug therapy				
Eczema or psoriasis affecting the auricle and auditor	y canal				
Allergy to components of air conduction hearing aids implants above	that are not included in one of t	he			
Stenotic canal severe enough to preclude use of an a	ir conduction hearing aid				
Excessive wax production that cannot be overcome was a series of the	-				
Excessive perspiration that cannot be overcome with	scheduled treatments				
d. Does the patient have middle ear disease?					
Additional Comments:					

\*Please fax completed form and medical records to 801-366-7449.