Clinical Policy: Cochlear Implant Replacements
Reference Number: CP.MP.14
Date of Last Revision: 07/22

See Important Reminder at the end of this policy for important regulatory and legal information.

Description
This policy outlines medical necessity criteria for the replacement of cochlear implants and/or cochlear implant components. The cochlear implant has 4 basic components: a microphone, worn externally behind the ear, which picks up sounds; an external speech processor which converts sounds to electrical signals; a transmitter and receiver/stimulator which forward the signals; and implanted electrodes, which stimulate the fibers of the auditory nerve.6

Policy/Criteria
I. It is the policy of health plans affiliated with Centene Corporation® that replacement of a cochlear implant(s) and/or its external components (external speech processor, controller, etc.) is considered medically necessary when any one of the following is present:
   A. The existing device(s) is no longer functional and cannot be repaired;
   B. A change in the member/enrollee's condition makes the existing unit(s) inadequate for the hearing-related activities of daily living and improvement is expected with a replacement unit(s);
   C. A sound processor replacement if the current processor is at least five years old.

II. It is the policy of health plans affiliated with Centene Corporation that replacement or upgrade of an existing, properly functioning cochlear implant and/or its external components (external speech processor, controller, etc.) is considered not medically necessary when requested only for convenience or to simply upgrade to a newer technology.

Background
Sensorineural hearing loss, or nerve deafness, is a type of hearing loss that results from problems with the inner ear, related to the cochlea, eighth nerve, internal auditory canal, or brain. A common cause of hearing loss in adults is presbycusis, a progressive condition caused by the loss of function of hair cells in the inner ear.7 Severe to profound hearing loss in children most often is caused by genetics, prenatal, perinatal, or postnatal causes.5 A cochlear implant, an electronic device surgically placed under the skin, bypasses the hair cells and directly transmits sounds through multiple electrodes, which stimulate the auditory nerve.7 Once the auditory nerve is activated, signals are sent to the brain. The brain learns to recognize these signals and the person experiences this as hearing.2

Cochlear implants have been studied since the 1950s and were approved by the FDA in adults in the mid-1980s.2,5 National Institute of Health (NIH) scientists determined cochlear implants to be cost beneficial.

Recent studies have been conducted evaluating the use of bilateral cochlear implants compared to unilateral implants. Many of these studies have shown that children obtained significantly higher hearing thresholds in the bilateral implants. Speech recognition scores in quiet and noisy
conditions were also improved in bilateral users. Studies also have shown better scores on sentence and word recognition tests for bilateral users.¹

Very little data has been published comparing differences between bilateral cochlear implants and cochlear implant with a hearing aid on the opposite ear. One small study showed improved localization abilities and speech perception scores for two former users of cochlear implant/hearing aid within the first 6 months after the second implant was activated. However, performance showed a slight decline after 6 months of use. Further studies are needed in this area to determine efficacy for bilateral cochlear implants in adults.¹

While evidence is increasing regarding the use of bilateral implants, bilateral implantation is not without problems. Limited nerve survival that remains may be asymmetrical, resulting in an unnatural pattern of neural activity in stimulation with electrical pulses. This asynchronous stimulation across devises might result in individual neural impulses which are unlikely to result in useful cues related to interaural differences. Also, bilateral implantation doubles the risks associated with surgical intervention and is very costly.²

**Coding Implications**
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<thead>
<tr>
<th>CPT®</th>
<th>Description</th>
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<tr>
<td>69949</td>
<td>Unlisted procedure, inner ear</td>
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<table>
<thead>
<tr>
<th>HCPCS Codes</th>
<th>Description</th>
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<tbody>
<tr>
<td>L8615</td>
<td>Headset/headpiece for use with cochlear implant device, replacement</td>
</tr>
<tr>
<td>L8616</td>
<td>Microphone for use with cochlear implant device, replacement</td>
</tr>
<tr>
<td>L8617</td>
<td>Transmitting coil for use with cochlear implant device, replacement</td>
</tr>
<tr>
<td>L8618</td>
<td>Transmitter cable for use with cochlear implant device, replacement</td>
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<tr>
<td>L6819</td>
<td>Cochlear implant, external speech processor and controller, integrated system, replacement</td>
</tr>
<tr>
<td>L8621</td>
<td>Zinc air battery for use with cochlear implant device and auditory osseointegrated sound processors, replacement</td>
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<tr>
<td>L8622</td>
<td>Alkaline battery for use with cochlear implant device, any size, replacement</td>
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<tr>
<td>L8623</td>
<td>Lithium ion battery for use with cochlear implant device speech processor, other than ear level, replacement</td>
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<tr>
<td>L8624</td>
<td>Lithium ion battery for use with cochlear implant device speech-processor, ear level replacement, each</td>
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HCPCS Codes | Description
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L8627 | Cochlear implant, external speech processor, component, replacement
L8628 | Cochlear implant, external controller component, replacement
L8629 | Transmitting coil and cable, integrated, for use with cochlear implant device, replacement

ICD-10-CM Diagnosis Codes that Support Coverage Criteria

| ICD 10 CM Code | Description |
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H90.3 | Sensorineural hearing loss, bilateral
H90.41 | Sensorineural hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side
H90.42 | Sensorineural hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side
H90.5 | Unspecified sensorineural hearing loss
Q85.00 | Neurofibromatosis, unspecified
Q85.02 | Neurofibromatosis, type 2
Z96.21 | Cochlear implant status

Reviews, Revisions, and Approvals

| Revision | Approval Date |
--- | ---
Initial approval date | 02/09
Added criteria for replacement of implants and components | 06/13 06/13
Clarified policy language to discuss replacement of implants or components only | 09/14 09/14
Background updated | 09/15 09/15
Converted into new template | 09/16 09/16
Clarified in II that replacements are not medically necessary when requested only for convenience/ to upgrade to newer technology. References reviewed and updated. | 09/17 09/17
References reviewed and updated. | 07/18 07/18
Added criteria for sound processor replacement if it is over 5 years old. | 10/18 10/18
Removed CPT 69717 and 69718 and replaced with CPT 69949 | 06/19
References reviewed and updated. Codes review. | 06/19 07/19
References reviewed and updated. | 06/20 07/20
Annual review. References reviewed and updated. Coding reviewed, added codes L8621 and L8622. Replaced all instance of “member” with “member/enrollee.” Changed “review date” in the header to “date of last revision” and “date” in the revision log header to “revision date.” Sent for specialist review. | 07/21 07/21
Annual review completed. Removed “or” in I.A. and I.B. Background updated with no impact to criteria. References reviewed and updated. | 07/22 07/22
References


Important Reminder
This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. “Health Plan” means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan’s affiliates, as applicable.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable Health Plan-level administrative policies and procedures.

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regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members/enrollees. This clinical policy is not intended to recommend treatment for members/enrollees. Members/enrollees should consult with their treating physician in connection with diagnosis and treatment decisions.

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Note: For Medicaid members/enrollees, when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

Note: For Medicare members/enrollees, to ensure consistency with the Medicare National Coverage Determinations (NCD) and Local Coverage Determinations (LCD), all applicable NCDs, LCDs, and Medicare Coverage Articles should be reviewed prior to applying the criteria set forth in this clinical policy. Refer to the CMS website at http://www.cms.gov for additional information.

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