

#### IV. その他の核遺伝子変異による ミトコンドリア病

ミトコンドリアに蛋白が輸送される時に、多くのサブユニットからなる輸送装置が使われる。外膜に結合する TOM 蛋白と内膜に結合している TIM 蛋白のうち、酵母の TIM8 蛋白に類似した DDP1 遺伝子に変異がある時に、ジストニアと難聴を呈する X 連鎖性の Mohr-Tranebjaerg 症候群が起きる<sup>8)</sup>。日本人でも報告がある<sup>24)</sup>。

別な X 連鎖性の疾患である Barth 症候群は、新生児や乳児期発症の拡張性心筋症、好中球減少症、ミオパチー、低身長、易感染性を呈する疾患であり、種々の電子伝達系酵素活性低下を伴う。責任遺伝子は G4.5, 別名 Tafazzin (TAZ) 遺伝子である<sup>25)</sup>。その機能はよく分かっていなかったが、最近になりリン脂質の合成やリモルディングに関わるアシル転移酵素との相同性が判明し、ミトコンドリア膜のリン脂質の研究が行われた。その結果、代表的なミトコンドリア膜のリン脂質であるカルジオリピン (CL) が患者では低下しており、さらに CL や他のリン脂質のリモルディングが障害されていることが示されている<sup>19)</sup>。これらリン脂質の異常とミトコンドリア異常との関連は今後の研究課題であるが、新しいミトコンドリア障害の機序として興味深い。

#### V. 今後の展望

ミトコンドリアの機能は、エネルギー産生だけにとどまらず、アポトーシス、カルシウムイオンの保管、活性酸素の発生など細胞機能にいろいろな影響を与える<sup>14)</sup>。これらの機能に関連して、パーキンソン病、アルツハイマー病、ハンチントン病などをはじめ、数多くの疾患と関連している。今後は、このような多くの疾患において、ミトコンドリアの持つどのような機能が病態と関係しているかという研究が重要であり、あらたな治療や予防に役立つ知見が得られる可能性が高いと考える。ミトコンドリアの研究は、新しい時代を迎えている。

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## Clinical Records

# Ototoxicity after use of neomycin eardrops is unrelated to A1555G point mutation in mitochondrial DNA

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### Abstract

Ear drops containing neomycin only rarely cause ototoxicity. The authors report on three patients with a tympanic membrane perforation who developed severe ototoxicity after use of eardrops containing 0.35 per cent neomycin. Mitochondrial DNA analysis revealed that there was no A1555G point mutation in these patients. This finding indicates that application of low concentration neomycin to the middle ear can cause severe inner ear damage even in humans who are not hyper-susceptible to aminoglycosides.

**Key words:** Neomycin; Cochlea; Hearing Loss, Sensorineural; Vertigo; Mutation; Drug Hypersensitivity

### Introduction

Neomycin is one of the most ototoxic drugs in modern medicine. Neomycin ototoxicity has been reported in all the routes of administration: oral, rectal, parenteral, intrapleural, wound irrigation, and aerosol spray. Neomycin primarily causes cochlear damage, especially when renal and gastrointestinal functions are altered.<sup>1</sup> Hearing impairment may develop, or continue to progress after systemic or topical application is discontinued.<sup>1,2</sup> Kavanagh and McCabe<sup>1</sup> reviewed cases of neomycin ototoxicity and concluded as follows: 'the usual latency is from two to six weeks after the onset of neomycin therapy. Since many patients received short courses of the drug, the hearing loss was often noticed days to weeks after the drug was discontinued. The onset of deafness may be severe and sudden but is usually progressive and may progress from six to 10 months, leaving the patient with a profound hearing loss.'

Several commonly used eardrops, including Cortisporin®, contain neomycin. When applied to the middle-ear space of guinea pigs, neomycin readily entered the perilymph in the cochlea through the round membrane<sup>3</sup> and at a concentration of 5 mg/ml or more caused dose-dependent ototoxicity.<sup>4,5</sup> Despite these experimental findings, hearing loss, vertigo, or both have rarely been reported after use of a neomycin otic suspension. Several large reference works failed to find such cases in humans (reviewed by Lind and Kristiansen<sup>6</sup>). An inquiry made by Swiss otolaryngologists showed that inner ear damage caused by antibiotic eardrops with ototoxic properties is estimated to occur in only one of 1 000 to 3 000 treatments.<sup>7</sup> Merfield *et al.*<sup>8</sup> found no

increase in the bone conduction thresholds in 44 children with chronic suppurative otitis media treated with five different topical aminoglycosides, four of which contained neomycin at a concentration of 0.33 or 0.35 per cent. Lind and Kristiansen,<sup>6</sup> however, reported that a patient with a ventilation tube to the middle ear developed severe hearing loss and vertigo after the use of eardrops containing 5 mg/ml neomycin. These findings suggest that humans may be less susceptible to neomycin applied topically to middle-ear spaces than are guinea pigs. Neomycin at the concentration of 0.35 per cent or less applied to middle-ear spaces may not be sufficient to cause inner ear damage in humans.

In Japan, Rindelon A® ear drops (Shionogi Co. Ltd.), which contain 0.35 per cent neomycin sulphate and 0.1 per cent betamethasone sodium phosphate in a 5 ml aqueous solution, are very commonly used. Approximately 19 000 000 bottles of these eardrops are sold annually in Japan. Despite the low concentration of neomycin, otologists know that this preparation may cause, albeit very rarely, inner ear damage. Why eardrops with low concentration neomycin are ototoxic in a limited number of cases is not known. The easy access of a neomycin otic suspension to the round window membrane in the absence of inflammatory oedema and secretion in the middle ear is considered a significant factor in producing this side effect.<sup>6</sup> Another possibility is that patients who experience inner ear damage after the use of a neomycin otic suspension may have hyper-susceptibility to aminoglycosides. Persons with an A-to-G transition at nucleotide (nt) 1555 of the 12S ribosomal RNA gene in the mitochondrial DNA (mtDNA) are reported to be very

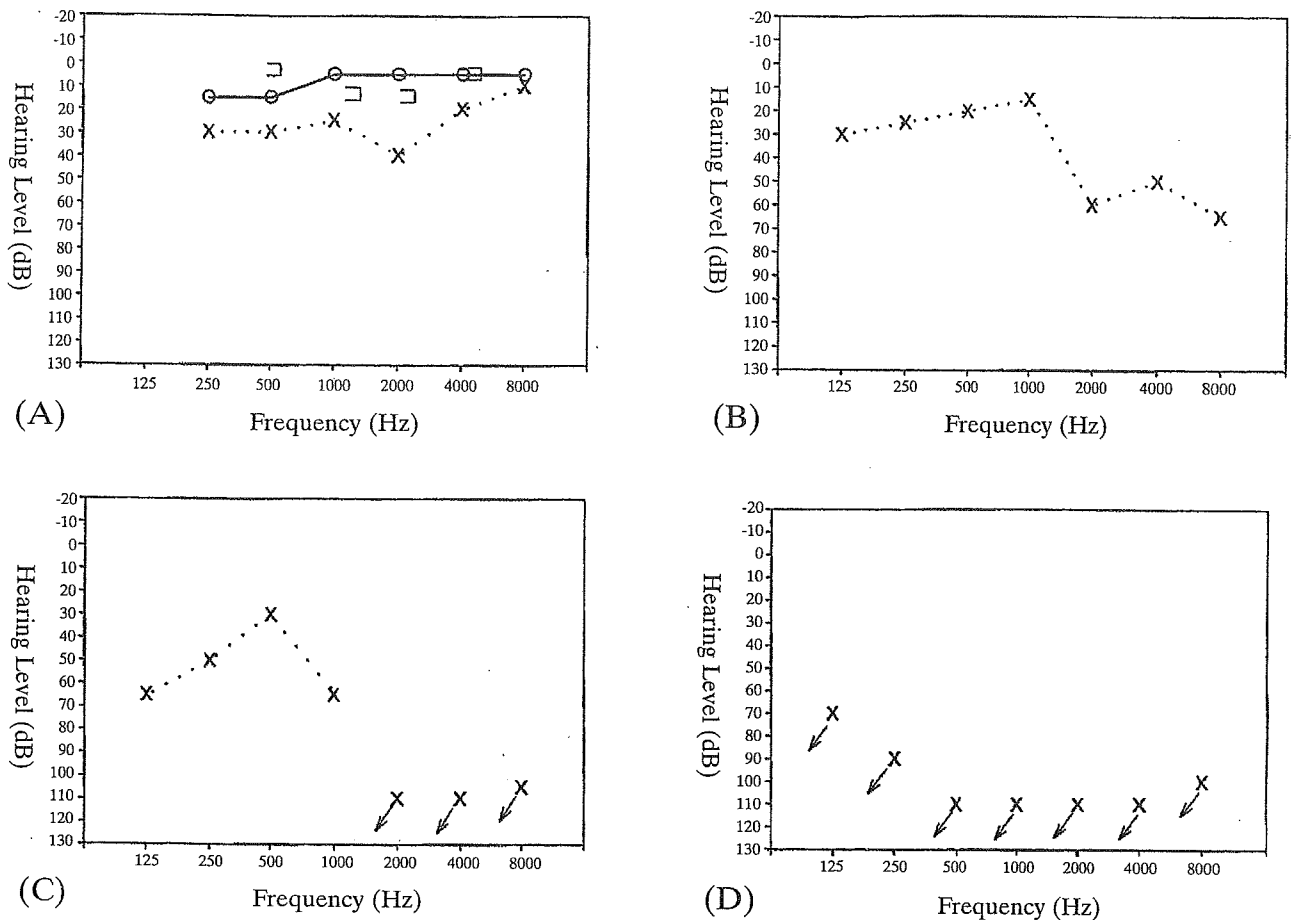


FIG. 1

Pure-tone audiograms for *Case 1*. (a) Slight conductive hearing loss on the left before use of Rindelon A® eardrops. (b) Sensorineural hearing loss at higher frequencies after the use of four bottles of the ear drops. (c) Progression of sensorineural hearing loss at all frequencies three months after cessation of eardrops use. (d) Complete deafness nine months after the use of Rindelon® eardrops.

susceptible to aminoglycosides.<sup>9-12</sup> The authors therefore investigated whether ototoxicity caused by a neomycin otic suspension was closely related to the presence of the A1555G mutation.

### Case reports

#### Case 1

A 50-year-old female suffering from tinnitus and mild hearing loss in the left ear, which had appeared one month before, visited the authors' clinic. She had been given six bottles of Rindelon A® ear drops over a three-month period at a private ENT clinic because of a slight serous discharge from her left ear. A pure-tone audiometric examination made before application of Rindelon A® ear drops showed mild conductive hearing loss on the left (Figure 1(a)). An otoscopic examination found a small perforation in the inferior part of the left tympanic membrane through which the mucoperiostium on the promontory looked intact. No discharge was present in the middle or external ear. A pure-tone audiometric examination showed an increase in hearing thresholds on the left at high frequencies (Figure 1(b)). Neuro-otological examinations found no abnormalities. An electronystagmogram (ENG) recorded no spontaneous, positional, or positioning systems. A caloric testing with

5 ml cold (4°C) saline produced nystagmus with a slow component velocity of approximately 35°/s in both ears. Magnetic resonance imaging (MRI) with Gd-DTPA enhancement excluded intracranial lesions such as a cerebellopontine angle (CPA) tumour. Although use of these eardrops was stopped, over the next six months her left-side hearing gradually worsened (Figures 1(c), 1(d)). Oral administrations of coenzyme Q, ATP and/or vitamin B12 were not effective. Nine months later, she was completely deaf in the involved ear.

#### Case 2

A 54-year-old male had been treated for one year for otitis media with effusion in the left ear at a private ENT clinic and had had a ventilation tube inserted four months before visiting this clinic. Before the insertion, he had mild conductive hearing loss on the left (Figure 2(a)). He had been given a total of four bottles of Rindelon A® ear drops because of occasional ear discharge through the ventilation tube. An otoscopic examination found that the ventilation tube had been placed in the antero-inferior quadrant of the left tympanic membrane, through which the mucoperiostium in the middle ear looked intact, and that there was no discharge. A pure-tone audiometric examination showed profound hearing loss on the left (Figure 2(b)). Neuro-otological examination found no

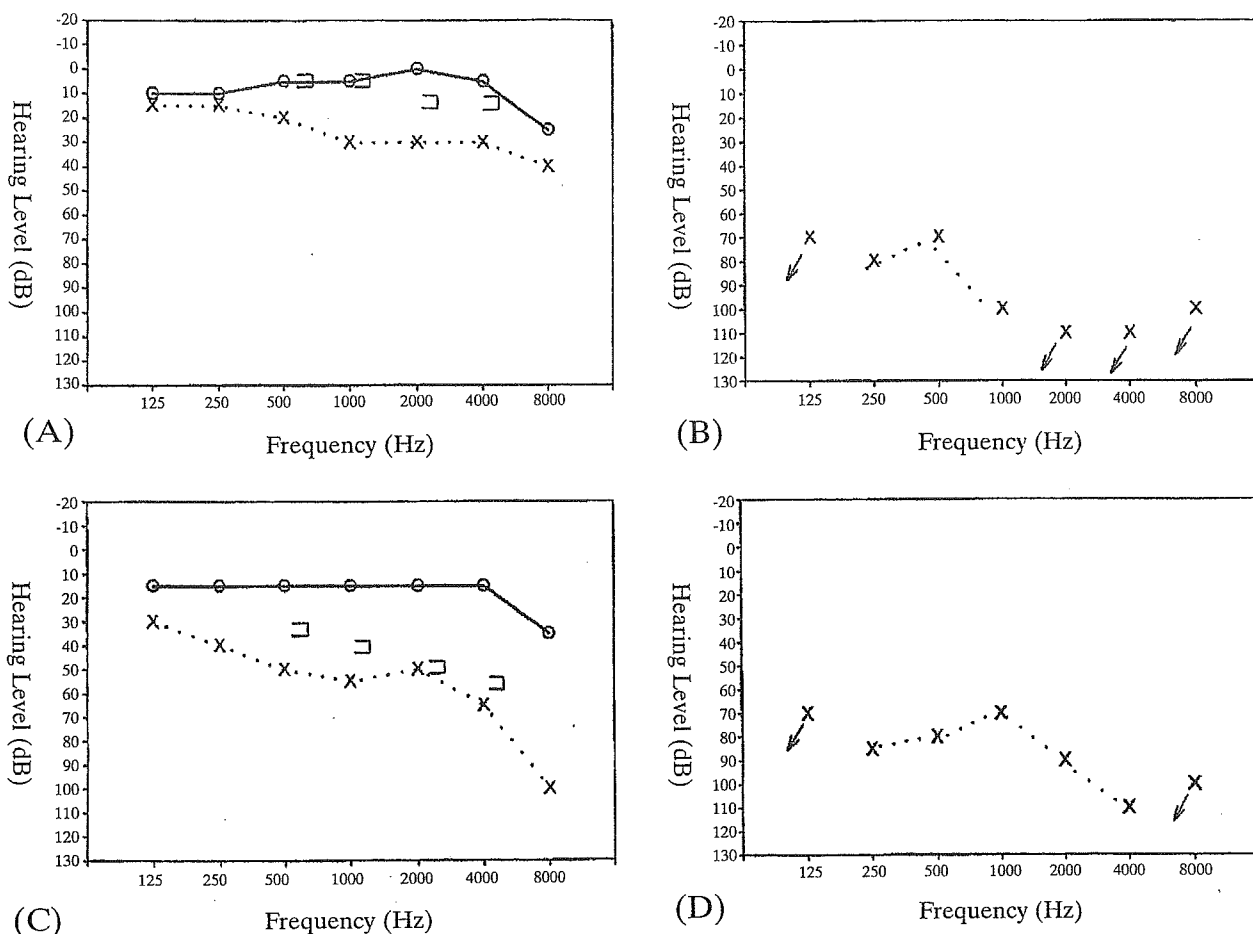


FIG. 2

(a) Mild conductive hearing loss on the left before the use of Rindelón A® ear drops in Case 2. (b) Profound mixed hearing loss on the left in Case 2 after the use of Rindelón A®. (c) Moderate mixed hearing loss on the left in Case 3 after the use of Rindelón A® ear drops. (d) Deterioration of hearing in Case 3 seen four months after the use of the ear drops.

abnormalities. An ENG recorded no spontaneous, positional or positioning nystagmus. A caloric testing with 5 ml cold (4°C) saline produced nystagmus with a slow component velocity of approximately 35°/s and 40°/s on stimulation of the right and left ear, respectively. Brain MRI with Gd-DTPA enhancement showed no abnormalities. The use of these eardrops was stopped, but when he revisited 12 months later, he was found to be completely deaf in the involved ear.

#### Case 3

A 58-year-old female had been given several bottles of Rindelón A® ear drops at a private ENT clinic six months before visiting this clinic for a serous discharge from her left ear. She experienced tinnitus and hearing loss in the left ear one month after using the eardrops. She had been given additional bottles of these eardrops one month before visiting this clinic, and two weeks after starting the use of these eardrops severe unsteadiness occurred. An otoscopic examination found a small perforation in the postero-superior quadrant of the left tympanic membrane, through which the mucoperiostium in the middle ear appeared to be intact. There was no discharge in the middle or external ear. A pure-tone audiometric examination showed moderate mixed hearing loss on the left (Figure 2(c)). Neuro-otological examinations revealed spontaneous horizontal-rotatory nystagmus, strongly

beating to the right. Caloric testing with 5 ml cold (4°C) saline produced nystagmus with a slow component velocity of approximately 30°/s on stimulation of the right ear, but no nystagmus was elicited on stimulation of the left ear with 20 ml cold saline. MRI with Gd-DTPA enhancement showed that there were no intracranial lesions such as a CPA tumour. Despite oral administration of a corticosteroid, ATP, and vitamin B12, spontaneous nystagmus continued over the next four months, and her left-side hearing deteriorated (Figure 2(d)).

#### Examination for the presence of A1555G point mutation in mtDNA

None of the three patients had close relatives with deafness, but their histories suggested that they might have been hyper-susceptible to aminoglycosides. The authors therefore examined whether any of the patients had an A1555G point mutation in their mtDNAs. The method used has been reported elsewhere.<sup>11</sup> In brief, total DNA was isolated from the patients' peripheral leukocytes. To determine whether the A1555G mutation was present, the mtDNA fragment surrounding the 12S rRNA mutation site was amplified by a polymerase chain reaction (PCR) that used the sense primer 5'-AAACTCAAAGGACCTGGCGG (nt 1160-1179) and antisense primer 5'-CGTCCAAGTGCAGTTTCCAG (nt 1598-1579). The fragments obtained were subcloned

into a plasmid vector by use of a TA cloning kit (Invitrogen, San Diego, Calif., USA) then directly sequenced in an ABI DNA sequencer. In the dot-blot hybridization, 2 µl of each amplified DNA was spotted onto nylon filters. A probe of 19-mer oligonucleotides five'-TACGACTTGCCTCCTCTAT (nt 1564-1545) was synthesized and labelled with  $\gamma$ -<sup>32</sup>P-ATP. The filter was hybridized with the labelled probe in 3 mol tetramethylammonium chloride (TMAC) solution for 3-5 h at 55°C. Then the filter was washed twice in rinse buffer at room temperature and twice more in TMAC solution at 58°C. The filter was exposed to Kodak XOMAT film for 1-5 h at -70°C.

Using this method, no abnormal sequences around nt 1555 were found in any of the patients. Moreover, none had the A1555G point mutation.

### Discussion

All patients in the present study had a tympanic membrane perforation unaccompanied by inflammatory oedema or discharge in the middle ear. They had received several bottles of Rindelon A® ear drops containing 0.35 per cent neomycin sulphate and 0.1 per cent betamethasone sodium phosphate. Patient 1 noticed hearing loss two months after using Rindelon A® ear drops and her hearing loss progressed nine months after the termination of the eardrop usage. Patient 2 developed profound hearing loss after use of Rindelon A® ear drops over four months and was found to become completely deaf in the involved ear when he revisited 12 months later. Patient 3 noticed hearing loss one month after using Rindelon A® ear drops. She developed moderate mixed hearing loss, which had progressed four months after the termination of the eardrop usage. By means of MRI, the presence of intracranial lesions such as CPA tumour was discounted in all patients. The latency from the onset of therapy and period of progression of hearing loss in these patients are similar to those reported in cases of neomycin ototoxicity.<sup>1,2</sup> The close time relationship between medication and the symptoms and the absence of intracranial lesions strongly suggest that the progressive hearing loss in these patients occurred due to the neomycin included in Rindelon A® ear drops that were applied to the perforated eardrum.

The concentration of neomycin in Rindelon A® eardrops is 0.35 per cent, the same as in Cortisporin® and Pediotic® otic suspensions commonly used in the USA. Despite its low concentration of neomycin, Rindelon A® ear drops induced severe inner ear damage in these patients. Following cessation of the use of this product, all these patients showed gradual deterioration of hearing. Patients 1 and 2 finally became completely deaf in the involved ear. In the involved ear of Case 3, vestibular function was virtually abolished. These cases of severe inner ear damage caused by eardrops containing a low concentration of neomycin raised the question whether the patients had hyper-susceptibility to aminoglycosides.

A homoplasmic A1555G point mutation in mtDNA is reported to be associated with maternally inherited aminoglycoside-induced deafness.<sup>9-12</sup> It has been shown using a rho<sup>0</sup> cell system that cells carrying this mutation had very high susceptibility to streptomycin.<sup>13</sup> Even very small amounts of aminoglycoside are reported to cause severe inner ear damage in persons with this mutation.<sup>10-12</sup> Patients with aminoglycoside-induced deafness and those with deafness of middle-age onset unrelated to aminoglycosides usually coexist in families harbouring the A1555G mutation,<sup>9-11</sup> but sporadic cases in which there was no family history of deafness have been reported.<sup>12</sup> Although none of the patients in this study had other

family members with deafness, the presence of these sporadic cases suggested that there might have been an A1555G point mutation. MtDNA analysis, however, showed that none of them had the A1555G point mutation. This shows that, even without the hyper-susceptibility to aminoglycosides caused by the A1555G point mutation, patients with a tympanic membrane perforation may develop severe inner ear damage after using eardrops containing neomycin.

- Previous studies have shown that aminoglycoside hypersensitivity is more severe in patients with a A1555G mutation in mitochondrial DNA
- In this paper three patients with tympanic membrane perforation developed ototoxicity following the use of neomycin based ear drops
- None of these patients had abnormalities of mitochondrial DNA
- These cases served to confirm the need for caution in the application of topical medication in all patients with safe chronic otitis media

No inflammatory oedema or pus was found in the middle-ear spaces of any of the patients. Eardrops are given usually when abundant pus and oedema are present in the middle ear of patients with a tympanic membrane perforation. A substance diluted by secretion therefore reaches the round window niche, which in such conditions is usually protected by thickening of the submucosal connective tissues. If there is no middle-ear inflammation, as in these patients, topically applied water-soluble substances can readily reach the round window membrane and gain access to inner ear fluids through that membrane. As previously reported,<sup>6</sup> easy access of neomycin solution to the round window membrane in the absence of inflammatory oedema and secretion may be a significant factor that contributes to the serious side-effects.

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# New Method of Using Reconstructed Images for Assessment of Patency of Intracochlear Spaces for Cochlear Implant Candidates

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**Objective/Hypothesis:** A new method of reconstruction using multidetector-row helical computed tomography (MDCT) was applied for assessment of cochlear patency in cochlear implantation candidates. **Study Design:** Descriptive study comparing radiologic and surgical findings in eight radiologically normal cochleas and four abnormal cases. **Methods:** In unusual cases including cochlear ossification and misinsertion of electrodes, intracochlear space was evaluated on two types of reconstructed images: the "basal turn plane" that was vertical to the modiolus and contained whole basal turn was effective in the assessment of the lining of the inner bony wall of the otic capsule; "cochlear cross plane" images allowed evaluation of the patency of the intracochlear space and the location of inserted electrodes at the sections vertical to the direction of insertion of the electrodes. **Results:** Preoperative evaluation was useful for predicting possible problems in electrode insertion and therefore helped in decision-making in regards to surgical strategies, especially in candidates with a high probability of cochlear ossification such as those with meningitis and otosclerosis. Furthermore, the new reconstruction method also provided useful information on the position of the cochlear implant electrodes after surgery. **Conclusions:** These unusual cases demonstrated that preoperative MDCT evaluation by our reconstruction method was useful for predicting possible problems in electrode insertion and decision-making in regards to surgical strategies, especially in cochlear implantation candidates with a high probability of cochlear stenosis. **Key Words:** Multidetector-row helical computed tomography, cochlear implantation, cochlear ossification, otosclerosis, meningitis.

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## INTRODUCTION

Partial or total obliteration of the cochlear scalae is frequently encountered in patients with deafness caused by otosclerosis, meningitis, and labyrinthitis.<sup>1</sup> Otosclerosis and meningitis can induce intracochlear changes that range from fibrosis to almost complete obliteration of the cochlea. In suppurative labyrinthitis, fibroblastic proliferation occurs within the perilymphatic spaces in the early period after bacterial penetration and is subsequently followed by ossification. In general, such ossification progresses from the base to the apex, resulting in complete labyrinthitis ossificans.<sup>2</sup>

Severe intracochlear obstructive changes have been found in a limited number of postlingually deaf adults but very commonly in children deafened by meningitis.<sup>3</sup> The House Ear Institute has reported that 64.0% of the 128 children who underwent cochlear implantation have become deaf because of meningitis.<sup>4</sup> In these cases, 21.3% of the cochleae showed radiologic evidence of ossified cochleas, but ossification of scala tympani was confirmed in as many as 33.9% during surgery.

Cochlear ossification evident on radiologic examination had previously been considered as a contraindication to cochlear implantation. The reasons for the contraindications included difficulty in insertion of the electrodes, increased resistance to spread of electrical stimuli, and possible damage to neural elements.<sup>1,5</sup> Since the description of alternative techniques such as the short insertion tunnel technique<sup>6</sup> and total drill-out technique,<sup>1,7</sup> several reports have demonstrated the safety and efficacy of cochlear implantation for patients with ossified cochleae.<sup>5,8,9</sup> The purpose of this report is to describe a new method using multidetector-row helical computed tomography (MDCT) to assess cochlear patency in cochlear implantation candidates, especially those with a high probability of cochlear ossification.

## MATERIALS AND METHODS

MDCT was performed before and after cochlear implantation, and reconstructed images were compared with the surgical findings. First, we applied our new method to eight patients who underwent cochlear implantation at our hospital. In both axial and coronal slices of high-resolution computed tomography

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(HRCT) and T2-weighted magnetic resonance images (MRI) taken before surgery, all showed no abnormal findings in the cochlea. The scala tympani was found during surgery to be widely open, and cochlear implant (CI) electrodes were inserted smoothly through a standard posterior tympanotomy approach in every case. Here, we describe the procedure used for the multiplanar reconstruction in a typical case. Figure 1 shows the preoperative images of a 9-year-old girl who had become deaf after mumps labyrinthitis 6 months earlier, and Figure 2 shows her images taken before and after insertion of the electrodes.

Axial projections of the temporal bone in both ears were taken in helical scan modes (0.5 mm slice width, 120 kV, 300 mA, pitch 3:1) with a multislice CT that has four detector rows (Toshiba Aquillon 16, New York, NY). The reconstruction spacing was 0.1 mm. The raw data were transferred to a workstation (Advantage Workstation version 4.0; GE Medical Systems, Milwaukee, WI), where three-dimensional (3D) reconstruction was performed. As shown in Figure 1, on the axial slice, in which the center of the modiolus was seen, we set a "basal turn plane" that was vertical to the modiolus and contained whole basal turn (Fig. 1A). Then, guidelines were set to pass through the modiolus on this basal turn plane (Fig. 1B). "Cochlear cross planes" (Fig. 2, A and B) were obtained so that they included both the modiolus and one of the guidelines. By rotating the guidelines around the axis, we could observe cochlear cross plane images consecutively. These reconstructed cochlear cross plane images contained the center of the fundus of the internal auditory meatus (IAM). Eighteen of these images taken every 10 degrees were printed on film. The presence of stenosis of intracochlear spaces caused by ossification or bony growth was evaluated on the basal turn plane images and consecutive cochlear cross plane images. As shown in Figure 1B, the lining of the inner bony wall of the otic capsule appeared smooth preoperatively in these patients on the basal turn plane images. Consecutive cochlear cross plane images showed lack of stenosis of the intracochlear spaces (Fig. 2, A and B). Figure 2, C and D, shows cochlear cross plane images taken after the insertion of 22 intracochlear electrodes and 2 mechanical support bands of the CI (Nucleus 24, Cochlear Ltd., Lane Cove, Australia). The image shown in Figure 2C depicts almost the same slice as the image shown in Figure 2A, and the image shown in Figure 2D corresponds with that shown in Figure 2B. These images clearly show the placement of the electrodes in the basal turn. These reconstructive procedures were performed cooperatively by radiologists and otologists, and a series of images for a subject was obtained in approximately 20 minutes.

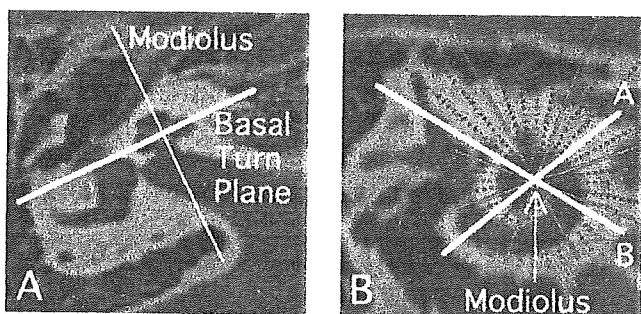


Fig. 1. (A) Preoperative axial slice of a 9-year-old girl in which the center of the modiolus is seen. A basal turn plane that was vertical to the modiolus and contains the whole basal turn is set. (B) The reconstructed basal turn plane image. On this image, the lining of the inner bony wall of the otic capsule is smooth. Guidelines are set to pass through the axis on this plane. Guideline "A" and guideline "B" indicate the cochlear cross planes demonstrated in Figure 2.

## CASE REPORTS

We describe four representative cases showing abnormal findings with our reconstruction method.

### Case 1

In a 62-year-old male with otosclerosis, axial and coronal HRCT demonstrated the presence of demineralization and destruction of the otic capsule on both sides, typical for the advanced and active phase of cochlear otosclerosis (Fig. 3A). However, it was difficult to determine whether there was space for electrode insertion. The basal turn plane images taken before surgery demonstrated irregular lining of the inner bony wall of the otic capsule in the right ear (Fig. 3B). Severe stenosis and lack of spaces in the lower portion of the basal turns were evident on the preoperative cochlear cross plane images (Fig. 4, A and B). These images also suggested the patency of mainly the scala vestibuli, which was located in the upper half of the basal turn. We explored the promontory in the right ear by the canal wall-up approach and found that the scala tympani around the round window niche were virtually occluded with ossified tissues. Using the transcanal drill-out technique,<sup>7</sup> we made a narrow tunnel, approximately 4 mm long, from the area just superioranterior to the round window niche to the ascending bend of the basal turn, where the patent scala vestibuli was exposed and a small amount of perilymph seepage was observed. Although the cavity of the scala vestibuli was narrow, 22 intracochlear electrodes and the 3 mechanical support bands of the CI (Nucleus 24) could be smoothly inserted into the scala vestibuli. The basal turn plane images (Fig. 3C) and cochlear cross plane images (Fig. 4, C and D) taken after surgery confirmed that the electrode array was fully inserted in the scala vestibuli of the basal turn.

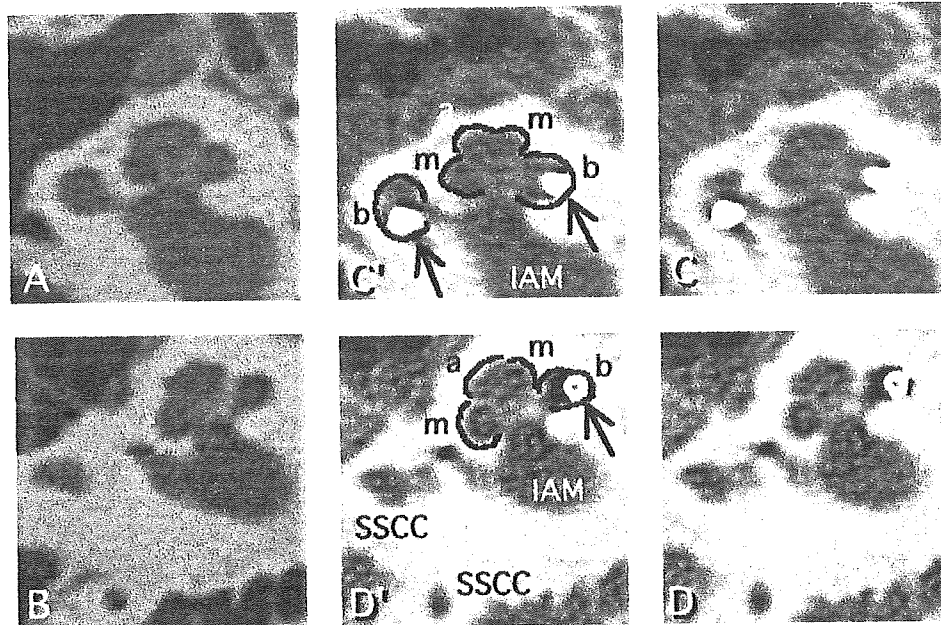
### Case 2

A 51-year-old male who had become deaf after meningitis underwent cochlear implantation at our hospital. Preoperative axial HRCT demonstrated chronic otitis media in the left ear and suggested a small lesion of ossification and slight stenosis in the basal turn of the right ear (Fig. 5A). The basal turn plane images taken before surgery also suggested this stenotic lesion (Fig. 5B). By axial and coronal HRCT, it was difficult to ascertain whether there was sufficient remaining space for electrode insertion. However, the cochlear cross plane images clearly demonstrated that the ossification in the basal turn was restricted to the portion near the modiolus and that there was remaining space outside the lesion (Fig. 5C). A total of 24 intracochlear electrodes of the CI (Medel C40+, MED-EL, Innsbruck, Australia) could be smoothly inserted into the scala tympani in the right ear through a routine posterior tympanotomy approach without additional technique. The cochlear cross plane images taken after surgery (Fig. 5D) confirmed that the electrode array was inserted in the remaining space, as observed in the preoperative images.

### Case 3

A 16-year-old male visited our hospital complaining of complete deafness. When he was 9 years old, he had become deaf because meningitis accompanying bilateral acute otitis media. He underwent cochlear implantation in the right ear at another university hospital 1 year later. The insertion of 22 intracochlear electrodes of the CI (Nucleus 22, Cochlear Ltd., Lane Cove, Australia) was successful, and he regained hearing. However, his hearing acuity began to deteriorate at age 15, and he was deaf again 3 months before his first visit to our hospital. At the first examination, the electrodes were found exposed in the external auditory canal. Preoperative axial HRCT demonstrated that the electrode array was completely drawn out from the cochlea, and soft tissue density mass was present around the round window

Fig. 2. Cochlear cross plane images of the same patient as demonstrated in Figure 1, taken before and after surgery. (A, B) On the preoperative cochlear cross plane images, note the patency of the intracochlear space of the basal, middle, and apical turns. (C, D) Cochlear cross plane images taken after the insertion of Nucleus 24 and their demonstrative schemes. Image C demonstrates almost the same slice as image A, and image D corresponds with image B. These images clearly show the position of the electrodes (arrows), which were placed in the basal turn (b). The apical turn (a), middle turn (m), internal auditory meatus (IAM), and superior semicircular canal (SSCC) are shown.



(Fig. 6A). To examine whether there was space for electrode reinsertion in the cochlea, we applied our new method of imaging. The basal turn plane images revealed ossification in the basal turn, although the lining of the inner bony wall of the otic capsule appeared smooth, except for the lesion. Severe stenosis and lack of spaces in the basal turns were evident on the cochlear cross plane images (Fig. 6C). Before surgery, we explained to the patient and his family that the reinsertion into the right cochlea might be impossible, and they consented that an option for the opposite side should be prepared for. We explored the tympanum in the right ear through a routine posterior tympanotomy approach and removed the electrode array. After removal of scar tissue in the tympanum, a fenestration into the cochlea was found at the point anterosuperior to the round window niche. Scar tissue was nearly totally removed, but because of the ossification in the intracochlear space, we failed to reinsert a sufficient number of electrodes. Another cochlea implant (Nucleus 24) was fully inserted in the left cochlea.

#### Case 4

A 58-year-old male was referred to our hospital from another hospital where he underwent cochlear implantation in the

left ear. He had become deaf because of meningitis 11 months earlier. According to the surgical record, a fenestration into the intracochlear space was successful, and perilymph seepage was observed. However, the operator found difficulty in inserting electrodes. Only seven intracochlear electrodes of the cochlea implant (Nucleus 24) were inserted. Because satisfactory recognition of speech was not obtained after the implantation, our method of cochlear imaging was requested to investigate the location of the electrodes. A postoperative basal turn plane image revealed that the tip of the array was not in the patent basal turn. The cochlear cross plane images suggested the electrodes were located outside the basal turn (Fig. 7B). 3D images also suggested that the tip had penetrated into the surrounding bony structure (Fig. 7D).

#### DISCUSSION

Our cases showed that preoperative evaluation of the intracochlear space with MDCT is useful for predicting possible problems in electrode insertion and therefore helps in decision-making in regards to surgical strategies, especially in cochlear implantation candidates with a high probability of cochlear stenosis. The smoothness of the

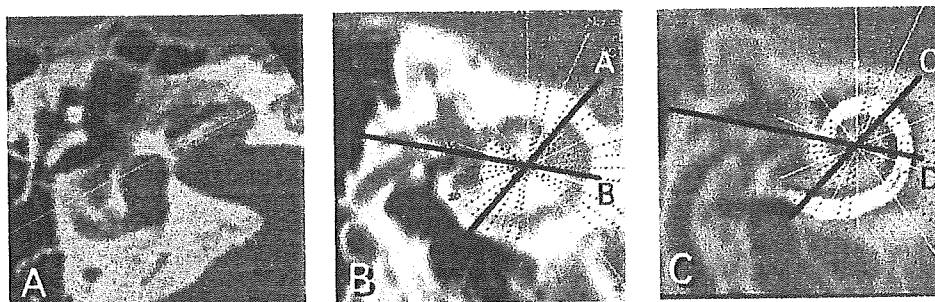


Fig. 3. (A) Conventional axial high-resolution computed tomographic (HRCT) image of a 62-year-old male with otosclerosis (case 1). Note the demineralization and destruction of the otic capsule, typical for the advanced and active phase of cochlear otosclerosis. (B) Preoperative basal turn plane image in case 1. Note the pericochlear lucency and irregular lining of the inner bony wall of the otic capsule. Guideline "A" and guideline "B" indicate the cochlear cross planes demonstrated in Figure 4. (C) Postoperative basal turn plane images taken after the insertion of Nucleus 24 in case 1. This image clearly demonstrates a fully inserted electrodes array in the basal turn. Guideline "C" and guideline "D" indicate the cochlear cross planes demonstrated in Figure 4.

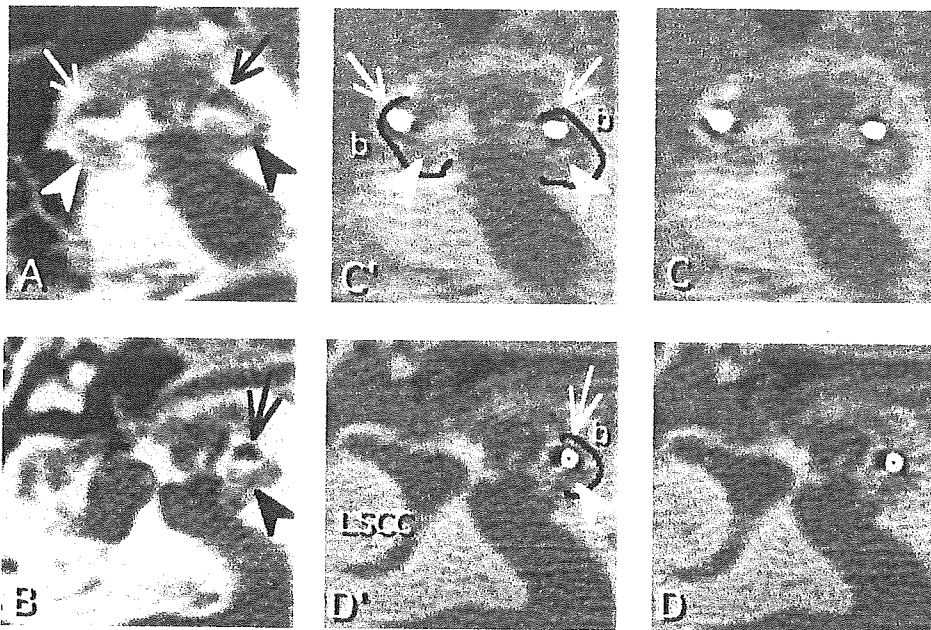


Fig. 4. Cochlear cross plane images taken before and after surgery in case 1. (A, B) Preoperative cochlear cross plane image demonstrates severe stenosis of the lower portion of the basal turns (arrowheads). In contrast, note the patency of the upper portion of the basal turn (arrows). (C, D) Cochlear cross plane images taken after the insertion of Nucleus 24 and their demonstrative schemes (C', D'). Image C demonstrates almost the same slice as image A, and image D corresponds with image B. These images clearly show the position of the electrodes (arrows), which are located in the upper portion of the basal turn (b).

inner wall of the otic capsule in the basal turn could be easily evaluated on the basal turn plane images. The cochlear cross plane images allowed evaluation of the patency of the intracochlear space at the sections vertical to the direction of insertion of the electrodes. In case 1, axial and coronal HRCT demonstrated severe demineralization and destruction of the otic capsule; however, the preoperative cochlear cross plane images demonstrated the area of patent spaces. This finding helped us to determine the most suitable surgical approach for such a difficult case. In case 2, a small lesion of ossification in the basal turn was suggested by axial HRCT, whereas the preoperative cochlear cross plane images showed the remaining space required for insertion of the electrode array. The patency

of the intracochlear space in the scala tympani was confirmed during the surgical procedure without additional drilling. Furthermore, the basal turn plane image and the cochlear cross plane image also provided useful information on the position of the CI electrodes after surgery. In case 3, the cochlear cross plane images provided excellent information on patent and ossified areas before reoperation. This information caused us to prepare for cochlea implantation in the opposite unoperated ear before we began the operation to remove the dislocated electrode array. In case 4, the misinsertion of electrodes was clearly recognized on the cochlear cross plane images and also on 3D images in a patient whose hearing acuity did not fully improve.

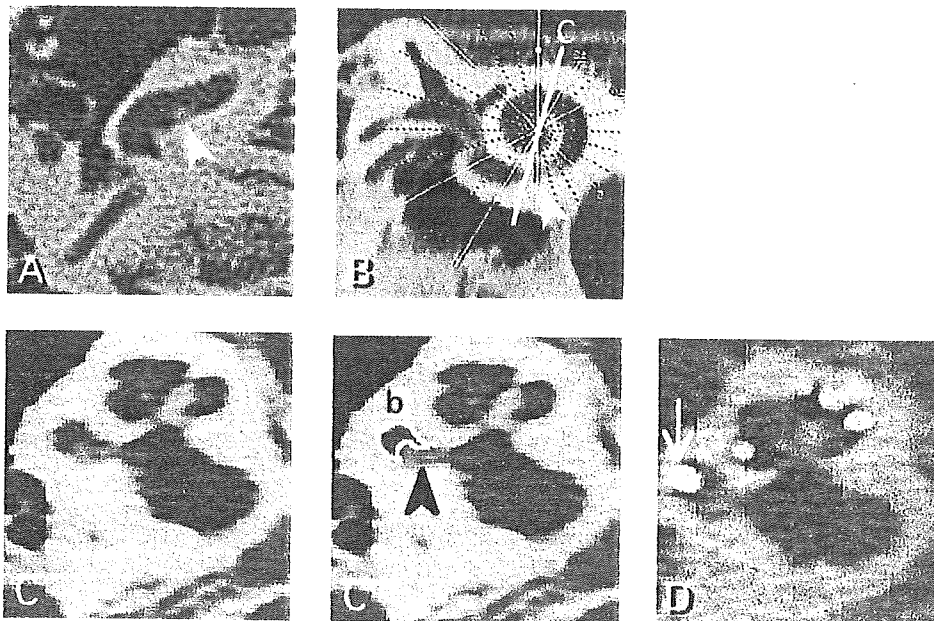


Fig. 5. (A) Axial high-resolution computed tomographic (HRCT) image of a 51-year-old male who had become deaf after meningitis (case 2). Preoperative HRCT suggested a small lesion of ossification (arrowhead) and slight stenosis in the basal turn of the right ear. (B) Preoperative basal turn plane image in case 2. In this image, the lining of the inner bony wall of the otic capsule is smooth except for a small lesion of ossification (arrowhead), which is sectioned on the cochlear cross planes (C) demonstrated in image C. (C) Preoperative cochlear cross plane image and its demonstrative scheme (C'). Ossification (arrowheads) near the modiolus and remaining space outside the lesion in the basal turn (b) are clearly demonstrated. (D) Cochlear cross plane image, which demonstrates almost the same slice as image C, was taken after the insertion of Medel C40+. This image confirms that the electrode array (arrow) was inserted in the remaining space identified in the preoperative image.

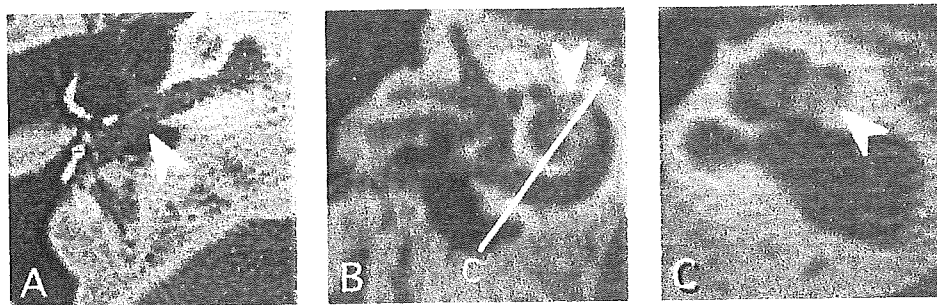


Fig. 6. (A) Axial high-resolution computed tomographic (HRCT) image of a 16-year-old male who had become deaf after meningitis accompanying bilateral acute otitis media (case 3). Preoperative HRCT demonstrated that the electrode array was completely drawn out from the cochlea, and soft tissue density was shown around the round window (arrowhead). (B) Preoperative basal turn plane image in case 3. Ossification in the basal turn (arrowhead) was revealed in this image, although the lining of the inner bony wall of the otic capsule appeared smooth, except for the lesion. Guideline "C" indicates the cochlear cross plane demonstrated in image C. (C) Severe stenosis and lack of spaces in the basal turn (arrowhead) was evident on the cochlear cross plane images.

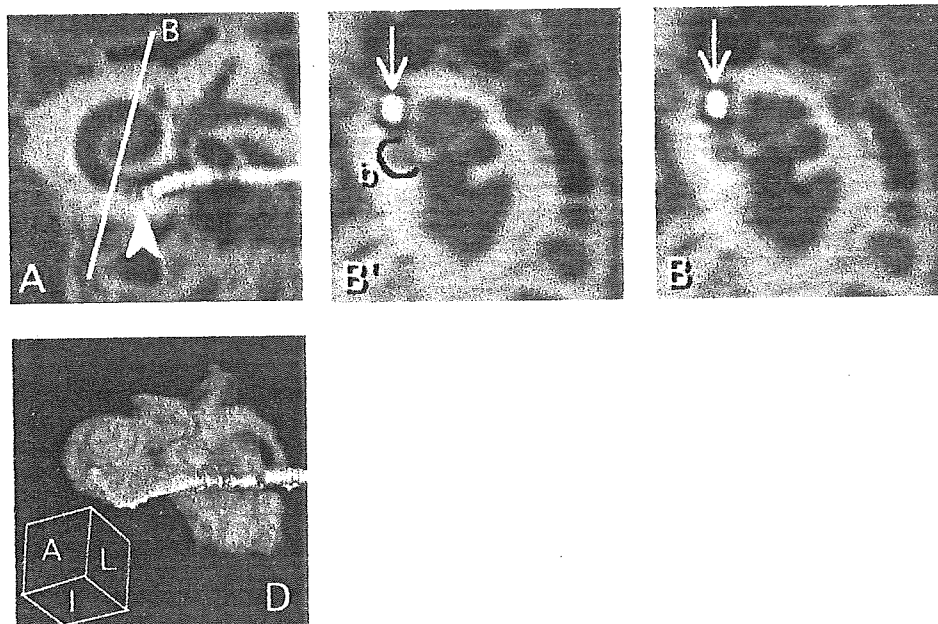
HRCT of the temporal bone has been considered useful for preoperative evaluation of candidates for cochlear implantation. Abnormal findings such as partial or complete cochlear ossification help to determine the most favorable side for implantation and allow surgeons to anticipate problems that may be encountered during device insertion. However, the two-dimensional (2D) axial images of HRCT cannot completely exclude the possibility of compromised cochlear patency. Small areas of ossification or soft-tissue obliteration in the cochlea, which can be an obstacle to smooth insertion of the electrode, may be missed.<sup>3</sup> Jackler et al.<sup>10</sup> reported a false-negative rate of 46% in their study using 2D axial images. Similarly, Seidman et al.<sup>11</sup> reported that in 32 cases of children with postmeningitis hearing loss, 2D scans with both axial and coronal slices had only a 53% accuracy in detecting intracochlear ossification. The resolution of the inner ear detail obtained by a CT scanner is limited by the width of slices and the partial volume effect. When there is a history of

meningitis or otosclerosis in a profoundly deaf person, these disadvantages are not acceptable.

High-resolution T2-weighted MRI of inner ears also provides valuable information of intracochlear space.<sup>12</sup> However, the assessment of detailed configuration of the lining of the inner bony wall of the otic capsule is preferable, especially when the additional drilling technique is expected to be used.

Recent advanced methods for the reconstruction of 3D images allow for significantly better visual recognition of fine structures in the temporal bone. Himi et al.<sup>13</sup> and Reisser et al.<sup>14</sup> have reported that the normal temporal bone structures are clearly recognized on helical CT images. Himi et al.<sup>13</sup> also obtained precise images of CI electrode routes similar to our 3D images in case 4 (Fig. 7D) by varying the angles of view. However, with the aid of an advanced workstation and software equipped with the recent standard device of MDCT, the combination of the basal turn plane images and the cochlear cross plane

Fig. 7. (A) Postoperative basal turn plane image of a 58-year-old male who regained no satisfactory hearing acuity after cochlear implantation with drilling (case 4). This image shows that the electrode array was penetrating into the bony wall (arrowhead) and that the tip of the array was not in the patent basal turn. Guideline "B" indicates the cochlear cross plane demonstrated in image B. (B) Cochlear cross plane images in case 4 and their demonstrative schemes (B'). These images clearly demonstrated that the electrodes (arrows) were located outside the basal turn (b). (D) Postoperative three-dimensional image of case 4. This image showed that the tip of the electrode array was penetrating into the surrounding bony structure. A, L, and I on the drawn cube indicate the anterior, lateral, and inferior direction, respectively.



images provided additional valuable information to evaluate cochlear patency of the intracochlear spaces. The MDCT protocol used in our study enabled excellent image quality on any reconstructed planes, and our method required neither special technical training nor additional cost for reconstructing the images. MDCT permitted routine use of very thin slices over large regions in short scanning time. The quality of MDCT images appeared to be superior, probably because of less motion artifacts because of the shorter scanning time. However, it was still difficult to discriminate between the scala tympani and the scala vestibuli by preoperative reconstructed images except in special cases, such as case 1.

## CONCLUSION

We describe a new reconstruction method using MDCT to assess cochlear patency in cochlear implantation candidates. The excellent results support the routine use of reconstructed MDCT images for evaluation of CI candidates.

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# Cortical Activation Shortly after Cochlear Implantation

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## Key Words

Cochlear implant · Positron emission tomography ·  
Plasticity · Word stimuli · Tone burst stimuli

## Abstract

We evaluated the cortical activations in postlingually deaf cochlear implant (CI) users in the early period (0–2 months) of CI usage. The subjects were 8 early CI users and 8 normal subjects. With tone burst stimuli (1 kHz) delivered to the right side, strong and broad activation of the ipsilateral (right) primary auditory cortex with 2 peaks and weaker activation of the contralateral (left) temporal lobe were observed in early CI users, in a clear contrast with the normal subjects in whom activation was observed in a small area of the contralateral (left) primary cortex. With word stimuli, activation of the superior frontomedian cortex presumably including the supplementary motor area and the neighboring cingulate gyri was observed in early CI users, which was absent in normal subjects. The activation in the immediate association cortices near the primary area was lower in early CI users, while the periphery of the association cortex seemed to be more mobilized.

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## Introduction

The cochlear implant (CI) is an artificial device which restores audition to profoundly deaf patients by directly stimulating the cochlear nerves. In CI users, auditory inputs are resupplied, suddenly at the first switch-on of the cochlear device, to the auditory pathways which are intact but have not been activated for a long period: a candidate condition for evaluating brain plasticity in audition. Positron emission tomography (PET) is a high-resolution imaging method to evaluate quantitatively the local blood flow which represents local synaptic activity. PET is the method of choice for studying CI users, because implanted patients cannot undergo functional MRI studies and because the localization power of electro-/magnetoencephalography remains far inferior to PET [Giraud et al., 2001]. Modified patterns of cortical activation have been reported in CI users with PET studies [Truy et al., 1995; Wong et al., 1999; Giraud et al., 2000; Naito et al., 2000]. The results of these studies, however, were not in perfect agreement, and most of them involved patients already rehabilitated for a relatively long period after implantation. Larger plasticity might be expected in an earlier period, since functional reorganization in the brain was shown in the early stage of idiopathic sudden sensorineural hearing loss, i.e. abrupt deprivation of the neural

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**Table 1.** Patient characteristics

Age/gender	Onset age of profound deafness right/left	Cause of deafness	CI usage before PET	Word score %
46/M	43/37	idiopathic (unknown)	1 month	80
58/M	57/57	idiopathic (unknown)	1 week	12
54/M	18/18	streptomycin toxicity	2 weeks	14
43/F	39/39	idiopathic (unknown)	2 weeks	22
45/F	43/25	idiopathic (unknown)	1.5 months	46
44/F	40/40	meningitis	2 weeks	64
53/F	52/52	otospongiosis	1 month	16
54/M	13/at birth	sudden deafness (cochlear aplasia on the left side)	2 months	6

input, an inverse condition to the present study [Po-Hung Li et al., 2003]. A small difference of cortical activations according to the length of use has been shown [Nishimura et al., 2000], but only in the long term (> 1 year of use). With early CI users, one report [Giraud et al., 2001] demonstrated smaller cross-modal plasticity in visual cortices in early CI users than in experienced users, and another [Giraud et al., 2001] showed smaller speech-related activation in the auditory cortex compared to rehabilitated users. In adult patients with lesions in the auditory cortices, such as due to infarction and epilepsy, possibly substitutive or compensatory activation of other cortical regions has been observed [Miura et al., 1998; Heiss et al., 1999; Warburton et al., 1999]. Brain plasticity in early CI users, if observed, may also contribute to clarify the mechanisms of such a possible substitution.

In the present study, we evaluated cortical activation in postlingually deaf patients implanted with CI devices by means of PET in an early period (0–2 months) of CI usage, when patterns of brain plasticity might be different from those already found in the later (>6–12 months) period.

## Methods

### Subjects

Eight normal male subjects (age 24–41 years, average 28.6 years) and 8 postlingually deaf patients implanted with cochlear implant devices (CI users: age 43–58 years, average 49.6 years), 4 males and 4 females, took part in the present study. The normal subjects were all right handed and had normal audition, without any history of psychiatric or neurological disorders. The CI users were also all right handed, were all implanted on the right side with a Nucleus Spectra 22 device but had variable speech intelligibility (single word recogni-

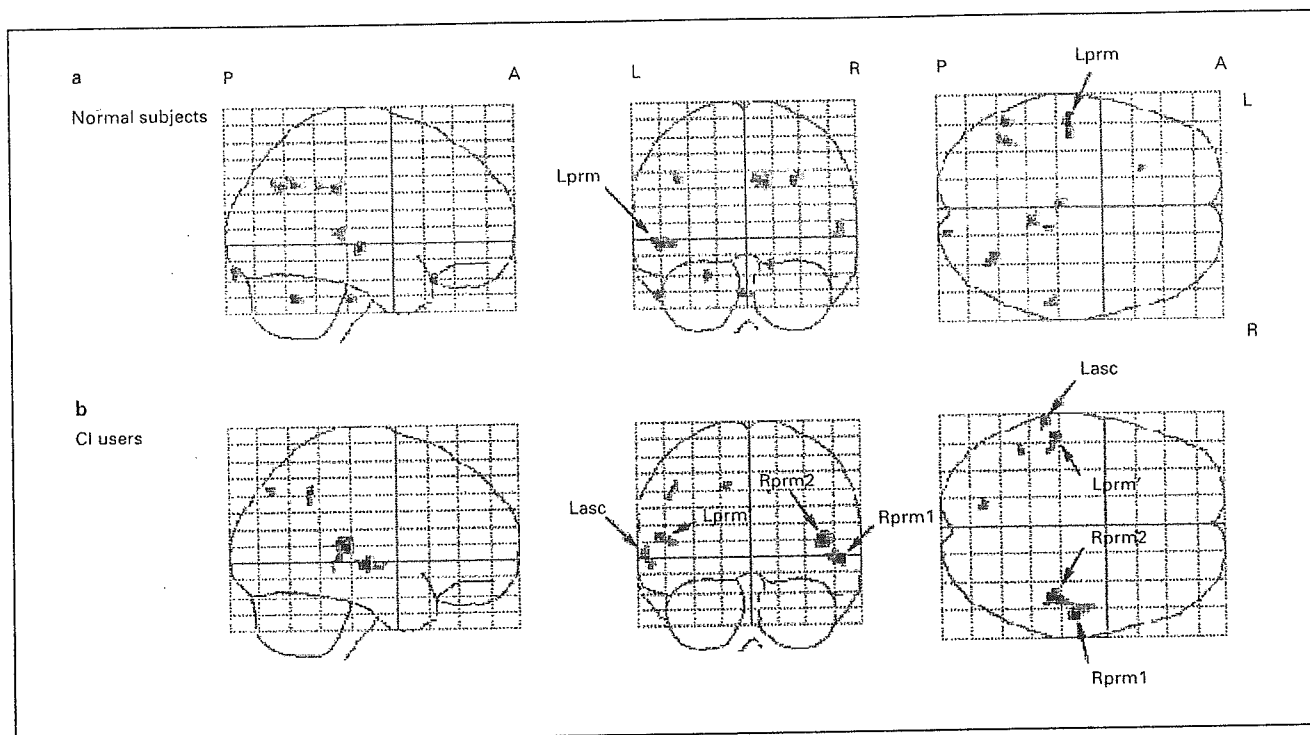
tion score: 6–80%, average 32.5%, without lip-reading) using the SPEAK strategy implemented in this device, in which natural spectral characteristics are more preserved, at the time of examination. They were relatively 'naive' users without much experience and rehabilitation with the device (1 week to 2 months after the switch-on of the device, median 24 days). Table 1 shows the patient characteristics. This study was performed in accordance with the guidelines of the Declaration of Helsinki and was approved by the ethical committee of the university hospital.

### Tasks and Stimuli

We used 3 conditions: silence, presentation of a train of 1-kHz tone bursts and word presentation. Each condition was repeated twice in a pseudo-random manner in which immediate repetition of the same condition was avoided. During the experiment, the subjects were instructed to close their eyes and silently listen to the sounds when the stimuli were presented, without repeating them in mind. Auditory stimuli were delivered to the right ear of the normal subjects via an ear phone, and to the right cochlea of the CI users via the CI devices. The left ears of the normal subjects were plugged with rubber plugs and covered by headphones, which attenuated the ambient noise by more than 35 dB. The intensity of the sound was adjusted to the so-called comfortable level, the term used in the CI fitting, in which the subject perceives the sound distinctly but does not find it too abrasive (corresponding to 60–70 dB) [Okazawa et al., 1996]. The tone burst stimuli consisted of 1-kHz tone bursts lasting 100 ms, repeated at a rate of 3 Hz. The word stimuli consisted of a series of 3-syllabled Japanese nouns presented at a rate of 1 Hz. The nature of the stimuli had been explained to the subjects, but the stimuli themselves were not presented beforehand, nor was the order of presentation of stimuli. The subjects were asked to report the nature of stimuli after each PET scanning. All the CI users as well as the normal subjects reported the right answers for all stimuli.

### PET Acquisition

Regional cerebral blood flow (CBF) was measured using the Headtome-IV system [Iida et al., 1989], following an intravenous bolus injection of  $H_2^{15}O$  (1480 MBq). Detailed procedures to acquire PET images appear elsewhere [Sakurai et al., 1996]. Briefly, the sound stimuli started 15 s before the scanning began and lasted for



**Fig. 1.** Projection images of statistical parametric maps, showing activation with tone burst stimuli in normal subjects (a) and CI users (b). The activated areas were thresholded at uncorrected  $p < 0.001$ , in order to show clearly the steep activation evoked by this pure-tone-like stimulus. P = Posterior; A = anterior; L = left; R = right. For abbreviations of brain areas, see text.

**Table 2.** Foci of maximal activation by tone burst stimuli within temporal lobes are shown in the order of Z values calculated by SPM99

	Area	Z value	Coordinates (x, y, z)
Normal subjects	Lprm	4.50	(-50, -20, 2)
CI users	Rprm1	4.42	(54, -20, -2)
		3.30	(50, -21, -4)
	Rprm2	4.36	(44, -32, 10)
	Lprm'	3.84	(-54, -30, 12)
	Lasc	3.84	(-62, -36, 0)

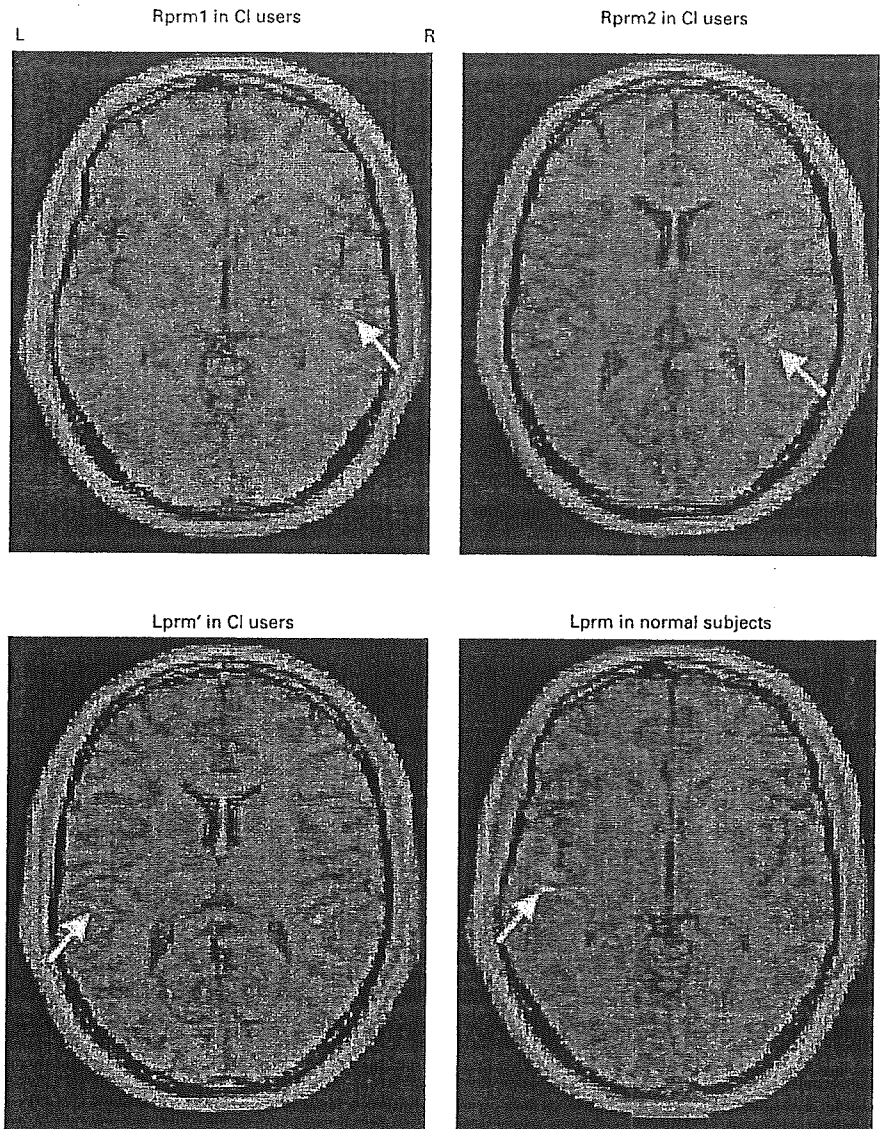
Lprm = Left primary auditory cortex (indicated as Lprm in normals and Lprm' in CI users); Rprm = right primary auditory cortex (Rprm1 and Rprm2: anterolateroinferior and posteromediosuperior portions of the right primary auditory cortex, respectively, in CI users); Lasc = left auditory association cortex. Coordinates (mm) are shown in the standard stereotaxic space of Talairach. Only the areas with peak Z values exceeding 3.3 are shown, i.e.  $p < 0.0005$ .

60 s, and the scanning lasted for 90 s. Fourteen slices with a pixel size of  $2 \times 2$  mm were obtained. The center-to-center slice distance was 6.5 mm. The CBF was measured based on Kety's single-compartment model. To compute the CBF precisely, arterial radioactivity was monitored with an on-line  $\beta+$  detector at the rate of 8 ml/min to provide an arterial time-radioactivity curve. The regional CBF values were normalized so that the whole CBF was constant (40 ml/100 g/min).

#### Data Analyses

Each set of 14 PET images was linearly interpolated into 53 slices (slice distance: 1.6 mm) and then converted to a format compatible with the SPM99 software (Statistical Parametric Mapping, realized by the Wellcome Department of Cognitive Neurology, Hammer-smith Hospital, London, UK) [Friston et al., 1991, 1994, 1995; Frackowiak et al., 1997], using the Igor Pro software (Wavemetrics Inc., USA). The standard pre-processings (realignment, normalization to the standard stereotaxic space [Talairach and Tournoux, 1988] and smoothing) and the statistical analyses were performed using SPM99 running on the Matlab 5.2.1 software (Mathworks Inc., USA) for the Windows 98 operating system (Microsoft Corp., USA). The positions of anterior and posterior commissures, necessary for these analyses, were determined by coregistration of PET images with MRI images of each subject, except 1 normal subject in whom





**Fig. 2.** Activated foci in the primary auditory area with tone burst stimuli. Each focus (Rprm1, Rprm2, and Lprm' in CI users and Lprm in normal subjects) determined by SPM99 are superimposed on the 'canonical' MRI images registered in SPM99.

MRI was unavailable. In this subject, the anterior-posterior commissure line was determined directly from the PET image. Statistical inference consisted of t tests at each voxel in the brain, which gave p and Z values for each voxel. A smaller p value or larger Z value represents stronger activation.

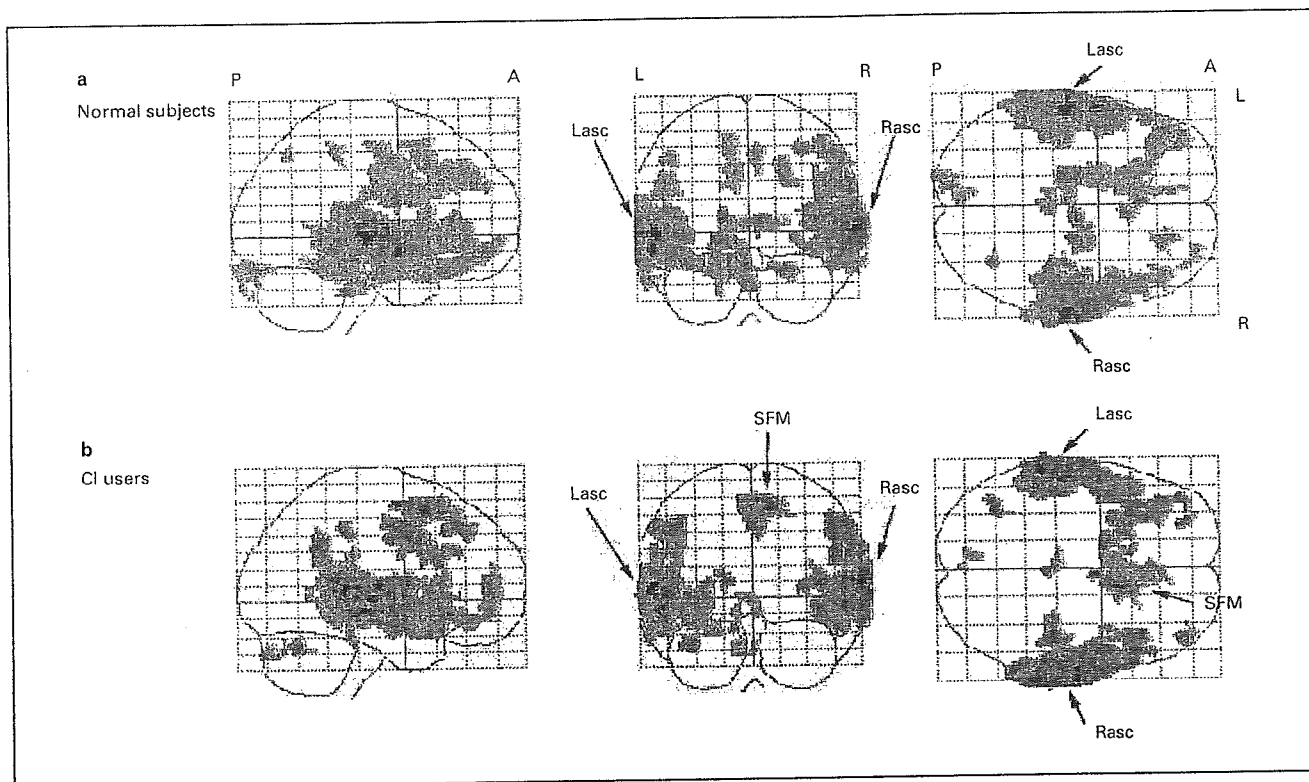
Activated brain areas were statistically determined by SPM99, which is based on statistical parametric maps using the general linear model. Both in normal subjects and CI users, the two conditions, silence and stimulation, were compared, and the brain areas activated by the stimuli were revealed. SPM is a statistical method developed to avoid the inevitable arbitrariness accompanied by the conventional region of interest method [Friston et al., 1995]. SPM can extract the essential activation related to a certain condition, eliminating the unimportant or noisy activation caused by individual dif-

ferences etc. We chose this method because of this advantage and because previous CI PET studies also employed this method, and did not discuss the activation case by case, which may permit numerous arbitrary interpretations.

## Results

### *Brain Activation by Tone Burst Stimuli*

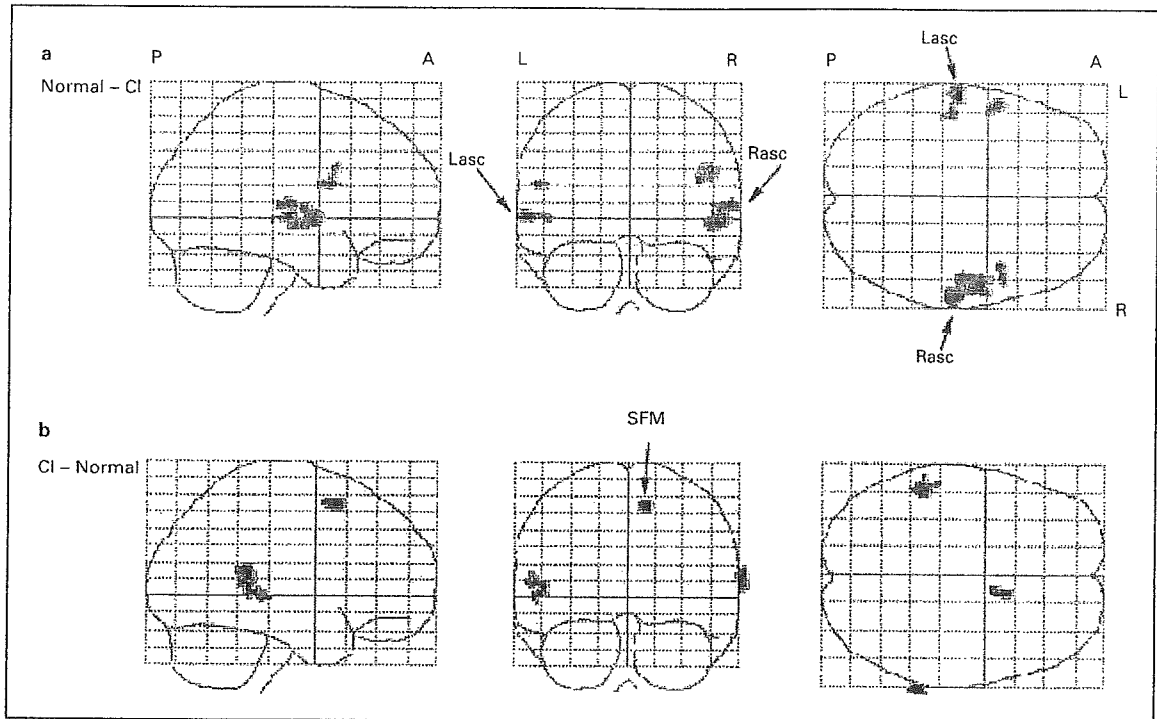
Tone burst stimuli (table 2) activated a small area in the left primary auditory cortex (Lprm, transverse temporal gyrus) in the temporal lobe of normal subjects (fig. 1a).



**Fig. 3.** Projection images of statistical parametric maps, showing activation with word stimuli in normal subjects (a) and CI users (b). The activated areas were thresholded at uncorrected  $p < 0.05$ , since a word stimulus evoked wide activation without very steep peaks. SFM = superior frontomedian cortex; for other abbreviations of brain areas, see text.

In CI users (fig. 1b), the pattern of activation was different. Strong and broad activation was observed in the right primary auditory cortex, ipsilateral to the sound stimuli. Two distinct peaks were found, one in the anterolateroinferior part (Rprm1) and the other in the posteromediosuperior part (Rprm2). Rprm1 had a symmetrical position to Lprm in normal subjects. Two foci with weaker activation than on the right side were observed in the left temporal lobe (Lprm' and Lasc). Lasc was situated in the left association auditory cortex (Lasc, superior or middle temporal gyri). Lprm' was located almost symmetrical to, but somewhat more lateral than, Rprm2 and corresponded to the border area between Brodmann's areas 41 (transverse gyrus) and 42 (superior gyrus) by reference to the standardized stereotaxic atlas [Talairach and Tournoux, 1988]. Figure 2 shows Rprm1, Rprm2 and Lprm' in CI users and Lprm in normal subjects, superimposed on the typical MRI image. All these 4 activated foci were located along and/or adjacent to the transverse temporal gyrus,

confirming activation in the primary auditory area because both cytoarchitectonic subdivision study [Brodmann, 1909] and functional study [Lauter et al., 1985] located the human primary auditory cortex on and adjacent to the transverse gyrus. However, Lprm' in CI users seems to involve the adjacent association area more strongly than the other 3. In reality, an activation in the temporal lobe has been considered to be in the primary cortex if it is near the transverse gyrus and is evoked by pure tones [Bilecen et al., 2000], because the activations of 2 distinct but adjacent areas, such as areas 41 and 42, cannot be distinguished on their border by the current resolution of imaging and because the morphological localization determined by gyri does not perfectly correspond to the cytoarchitectonic configuration (Brodmann's areas). Therefore, we regard Lprm' in CI users as an activation in the primary area, rather than an extraordinary plasticity occurring in the association cortex.



**Fig. 4.** Regions where activation with word stimuli was stronger in normal subjects than in CI users (a) and vice versa (b). The areas were thresholded at uncorrected  $p < 0.001$ .

#### Brain Activation by Word Stimuli

Word stimuli (table 3) activated bilateral auditory association cortices (left: Lasc, right: Rasc) in normal subjects (fig. 3a).

In the CI users (fig. 3b), in addition to bilateral auditory association cortices (Lasc and Rasc), the right superior frontomedian cortex (SFM) presumably including the supplementary motor area and the neighboring midline anterior cingulate gyri were also activated. In association cortices, the peak Z values in CI users were smaller than in normal subjects, and the statistical parametric map images gave an impression of broader activation (small activated foci scattered in the bilateral association cortices, in contrast to more compact, concentrated activated foci in normal subjects). However, since the activated foci in the temporal lobe largely overlap with word stimulation and the difference is not so easy to determine as with tone bursts, we investigated the differential activation between CI users and normal subjects (fig. 4). Activation in bilateral association cortices near primary areas, corresponding to Brodmann's area 42, was higher in normal subjects. In CI users, activation in the right SFM was

**Table 3.** Foci of maximal activation by word stimuli in 3 areas of interest are shown in the order of Z values calculated by SPM99

	Area	Z value	Coordinates (x, y, z)
Normal subjects	Lasc	5.52	(-58, -22, 2)
		4.96	(-58, 0, -10)
		3.90	(-68, -22, 6)
	Rasc	5.49	(66, -22, 2)
		5.02	(68, -14, 2)
		4.74	(60, -8, 4)
CI users	Lasc	5.01	(-60, -36, 6)
		4.51	(-54, -24, -6)
		4.19	(-50, -38, 4)
	Rasc	4.43	(62, -16, 10)
		4.32	(68, -40, 12)
		4.17	(72, -20, 0)
	SFM	4.30	(12, 12, 54)
4.20		(4, -4, 60)	

Lasc and Rasc = Left and right auditory association cortices, respectively; SFM = right superior frontomedian cortex. Coordinates (mm) are shown in the standard stereotaxic space of Talairach. Only the areas with peak Z values exceeding 3.3 are shown, i.e.  $p < 0.0005$ .

higher than in normal subjects. We also found that activation in CI users is higher at the posterior end of bilateral association cortices (posterior end of Brodmann's area 22).

## Discussion

### *Differences in Cortical Activation between Normal Subjects and CI Users* Tone Burst Stimuli

Cortical activation patterns by the tone burst stimuli showed an outstanding difference between normal subjects and CI users. Whereas a small area in the *contralateral* primary auditory cortex was activated in normal subjects, a much broader area in the *ipsilateral* primary cortex was predominantly activated, with 2 separate peaks, in early CI users. No comparable plasticity in the auditory primary cortex has been reported. The broadness of this ipsilateral primary cortex activation may be due to stimulation of a larger population of cochlear neurons, in contrast to a very limited stimulation in normal subjects, owing to inevitable current leaks of CI devices and/or may signify that such ipsilateral projection is new and therefore preliminary, failing to converge on a small central processing area. The functional auditory projection to the primary cortex was shown to shift from contralateral to bilateral after unilateral deafness, possibly due to disinhibition from the contralateral inputs [Vasama et al., 1995; Scheffner et al., 1998; Bilecen et al., 2000; Ponton et al., 2001], especially with left ear unilateral deafness [Khosla et al., 2003]. Although the contralateral activation remained dominant in such cases, predominant ipsilateral activation was found in the very acute phase of sudden deafness [Suzuki et al., 2002; Po-Hung Li et al., 2003]. In the present study, the ipsilateral projection (right ear to right primary cortex) in early CI users may partly be attributed to lack of inhibition from the contralateral (left) ear. However, since we tested only right-sided CI users, we dare not assert that the left-sided early CI users will show similar ipsilateral activation, since a difference in cortical activation according to the deafened side in patients with unilateral deafness was found [Khosla et al., 2003].

Our present results may not be compared directly to previous studies, since no previous PET studies in CI users employed tone burst or pure-tone stimuli. Electric and magnetic responses over the ipsilateral (right) hemisphere were observed in 1 patient implanted on the right side with tone burst stimuli [Pelizzone et al., 1986].

Giraud et al. [2001] found activation of the contralateral (right) primary cortex with a mixture of speech and noise stimuli in early left-sided CI users. One study with a brief square pulse, which corresponds to a click stimulus in normal-hearing subjects, showed that the activation was contralateral in 2 of the 3 subjects tested and was ipsilateral in 1 subject [Truy et al., 1995], although the length of the usage of devices was not noted. Thus, the results were variable depending on the side of stimulation, type of stimulus and period of rehabilitation. However, PET studies with postlingual CI users including long-term users, whose subjects were similar to ours, have shown that the white noise activated the contralateral primary cortex more strongly than the ipsilateral cortex [Naito et al., 1995; Okazawa et al., 1996]. Assuming that the difference between tone burst stimuli and white noise stimuli could be ignored, we hypothesize that the prominent right-sided ipsilaterality observed in the present study may diminish in the long-term use of CI devices, i.e. in the course of auditory rehabilitation.

In discussing the lateralization of cortical responses, it is important to note the large difference among different types of sound stimulation. With meaningful speech stimuli, bilateral temporal gyri (primary and association cortices) are strongly activated with higher activation on the left side [Scott et al., 2000]. With word stimuli, the pattern of activation is similar but less strong with less marked left-sidedness [Wise et al., 2001]. With pure-tone or tone burst stimuli, a small area of the contralateral temporal gyrus (primary cortex) is activated [Lauter et al., 1985]. Cortical activation with white noise or click stimuli, which has a wide frequency range, shows an intermediate and obscure pattern between speech and pure tone, i.e. bilateral (contralateral > ipsilateral), broad but weaker activation than that caused by speech stimuli [Naito et al., 1995]. Moreover, the significant activation of the primary cortex was not consistently shown (only small to no statistical significance) in previous PET studies with white-noise stimuli [Okazawa et al., 1996; Naito et al., 1997, 2000]. Lateralization differences of cortical activation according to gender have been controversial, although a recent large-scale report denied intersexual differences [Frost et al., 1999]. Hemispheric asymmetry affected by aging was also reported [Bellis et al., 2000]. However, these were shown only with speech stimuli and only the laterality, not the precise location of activated foci, was noted. Such differences with pure-tone stimuli were not reported and seem not plausible with bilaterally normal-hearing ears, considering the predominant contralateral connection of the primary cortex. In fact, exact age/gen-