Vestibular Implant Imaging

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ABSTRACT

SUMMARY: Analogous to hearing restoration via cochlear implants, vestibular function could be restored via vestibular implants that electrically stimulate vestibular nerve branches to encode head motion. This study presents the technical feasibility and first imaging results of CT for vestibular implants in 8 participants of the first-in-human Multichannel Vestibular Implant Early Feasibility Study. Imaging characteristics of 8 participants (3 men, 5 women; median age, 59.5 years; range, 51–66 years) implanted with a Multichannel Vestibular Implant System who underwent a postimplantation multislice CT (n = 2) or flat panel CT (n = 6) are reported. The device comprises 9 platinum electrodes inserted into the ampullae of the 3 semicircular canals and 1 reference electrode inserted in the common crus. Electrode insertion site, positions, length and angle of insertion, and number of artifacts were assessed. Individual electrode contacts were barely discernible in the 2 participants imaged using multislice CT. Electrode and osseous structures were detectable but blurred so that only 12 of the 18 stimulating electrode contacts could be individually identified. Flat panel CT could identify all 10 electrode contacts in all 6 participants. The median reference electrode electrode insertion depth angle was 9° (range, −57.5° to 45°), and the median reference electrode insertion length was 42 mm (range, −21–66 mm). Flat panel CT of vestibular implants produces higher-resolution images with fewer artifacts than multidetector row CT, allowing visualization of individual electrode contacts and quantification of their locations relative to vestibular semicircular canals and ampullae. As multichannel vestibular implant imaging improves, so will our understanding of the relationship between electrode placement and vestibular performance.

ABBREVIATIONS: FF = full-field; FPCT = flat panel CT; HR = high-resolution; MSCT = multislice CT; SCC = semicircular canal; MVI = Multichannel Vestibular Implant System

A. Although individuals with a unilateral vestibular deficit and 1 normal labyrinth usually compensate well via rehabilitation exercises and adaptation, those with bilateral vestibular hypofunction often have degraded visual acuity during head movement, postural instability, and chronic disequilibrium.1–3 When bilateral vestibular hypofunction results from ototoxic drug exposure, Menière disease, genetic defects, or other inner ear dysfunction sparing the vestibular nerve and central pathways, an implantable neuroelectronic prosthesis that measures 3D head rotation and stimulates the vestibular nerve with motion-modulated electrical pulse trains could substantially improve quality of life.4,5

Vestibular implants are similar to commercially available cochlear implants in that they include an external unit that powers and communicates with an implanted inner ear stimulator via a transcutaneous inductive link.4,5 The external unit includes a head-worn unit (for sensing head motion and delivering power and signals to the implanted stimulator) and a power and control unit containing a battery and microprocessor. As in cochlear implant systems, the head-worn unit and implanted stimulator each contain at least 1 magnet to hold the head-worn unit on the scalp over the implant. Unlike cochlear implant systems, vestibular implant systems sometimes include ≥1 additional magnet on each component to facilitate retention of the head-worn unit. Vestibular implant electrode arrays typically are much smaller than cochlear implant electrode arrays and are implanted in the semicircular canal (SCCs) near the ampullae, where the vestibular nerves branches terminate.4 Variations on this approach have included an

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extraluminal approach (in which the surgeon attempts to expose and place electrodes near vestibular nerve branches in the distal-most aspect of the internal auditory canal and singular nerve canal); implantation of electrode arrays near the utricle and saccule; simultaneous or delayed placement of a separate electrode array in the cochlea; and variations in location of the stimulation reference electrode, which can be implanted in the labyrinth or a subperiosteal pocket (as an alternative surgical procedure) or integrated with the stimulator housing.

Empiric studies in animals and finite element models of current flow in the labyrinth indicate that electrode distances to target and nontarget nerve branches are key determinants of the strength and selectivity of stimulation. Therefore, precise knowledge of electrode location, as provided by high-resolution postoperative imaging, can provide information helpful for both prognosis and guiding the choice of which electrodes to activate and which stimulus parameters to use. Postoperative imaging performed as a part of a vestibular implant operation is useful to confirm the location of implanted electrode arrays, measure the depth of insertion and electrode position relative to vestibular nerve branches, and detect kinking, damage, or displacement. MR imaging is unsuitable for assessing electrode location because of field interactions that distort images and can displace the magnet of the implant, poor air-bone contrast within the temporal bone, and the inability of MR imaging to directly image the platinum/iridium wires and silicone that make up electrode arrays (other than by imaging displacement of inner ear fluids). Multislice CT (MSCT) is the better technique for characterizing cochlear implant position and is the current de facto standard, given its greater spatial resolution and better contrast among bone, air, metal, and fluid.6,7 Metal artifacts, however, can significantly degrade image quality. Flat panel CT (FPCT), a relatively new imaging technique that yields tomographic reconstructions from images acquired using a C-arm x-ray system with flat panel image detectors, provides excellent visualization of high-contrast structures with better spatial resolution than MSCT.8,9 In particular, FPCT produces images with sufficient resolution to precisely quantify cochlear electrode contact locations in a clinical setting.6,7,9,10

The purpose of this study was to present the very first imaging results of CT in vestibular implant imaging. In this report, we present the technical aspects and imaging performance of MSCT and FPCT in assessing the intravestibular position of implanted electrode arrays in 8 participants in the first-in-human Multichannel Vestibular Implant Early Feasibility Study (clinicaltrials.gov, NCT02725463), and we suggest important features that should be reported in post-vestibular implantation imaging studies as well as a protocol for FPCT imaging of vestibular implants.11

MATERIALS AND METHODS

This study was conducted under a protocol approved by the Johns Hopkins institutional review board (No. NA_00051349) and was registered on the clinicaltrials.gov data base (NCT02725463).

Vestibular Implant Electrode Array Design and Implantation

The implanted stimulator component of the Multichannel Vestibular Implant System (MVI; Labyrinth Devices) is a CONCERTO cochlear implant stimulator (MED-EL), modified for implantation in the SCCs (Fig 1A). It includes an electrode array with stimulation electrodes, a stimulation return and recording reference electrode, hermetically encapsulated electronics, 3 magnets, and an antenna coil for transcutaneous inductive transmission of power and control signals that the implant receives from the external system component.

The MVI is implanted via a postauricular incision and mastoid approach similar to that typically used for cochlear implantation or labyrinthectomy, except that no entry is made into the cochlea and the SCCs are identified but not destroyed. Instead openings are made into the labyrinth for electrode array insertion (in the superior SCC ampulla, horizontal SCC ampulla, posterior SCC thin segment, and near the common crus). The electrode array (Fig 1B, C, and D) consists of a silicone carrier and comprises 10 platinum/iridium electrodes: 2 linear arrays of 3 electrodes each, joined to form a forked array inserted into the horizontal and superior ampullae; 3 on a linear array implanted in the posterior canal; and a braided platinum/iridium wire reference/return electrode inserted either into the common crus or in a subperiosteal pocket outside the temporal bone. Electrodes are spaced 0.2 mm apart in the silicone carrier for the forked array for the horizontal and

FIG 1. A, The MVI stimulator comprises 3 fixation magnets, an inductive coil link, electrical current stimulator circuitry, a stimulation electrode array, a stimulation reference electrode, and a recording reference electrode. The electrode array includes a 3-electrode shank for the posterior canal (E5–E7), a forked subarray with 2 shanks for the horizontal (E6–E8) and anterior (E9–E11) canals, and a stimulation reference electrode (D). eCAP indicates electrically evoked compound action potential. Reprinted with permission from Labyrinth Devices, LLC, 2019.
superior ampullae and 0.3 mm apart in the silicone carrier for the posterior SCC.

Participants
Eight participants (3 men, 5 women; median age, 59.5 years, range, 51–66 years) disabled by bilateral vestibular hypofunction were implanted unilaterally with the implanted receiver/stimulator of the MVI. Three participants were implanted in the right ear, and 5 participants, in the left ear. Table 1 summarizes demographic information.

Image Acquisition
After implantation, all subjects were scanned with either an MSCT (Somatom Sensation; Siemens) or a C-arm-based FPCT platform (Artis zee biplane; Siemens). MSCT was performed using standard clinical imaging parameters for temporal bone CT, with orientation of “axial” slices pitched to align with a plane through the horizontal SCC. Scanning was performed with 0.6-mm collimation, 120 kV, and 320 mAs. FPCT (DynaCT; Siemens) evaluation was performed using a flat panel angiography system (Axiom Artis zee; Siemens) and commercially available software (syngo DynaCT; Siemens). The participant was placed supine on the angiography table, and the head was taped in place to limit participant motion. When we prepared the DynaCT acquisition, attention was paid to collimate the VOI to include only the temporal bones (craniocaudal collimation from just above the petrous ridges to just below the mastoid tip). A 20-second FPCT acquisition of the head was performed using the following parameters: 109 kV, small focus, 200° rotation angle, and angulation step of 0.4° per frame. FPCT was performed in 2 modes: full-field (FF) and high-resolution (HR). The FF mode uses top and bottom collimation, whereas the HR mode has collimation in all planes, allowing focal acquisition of the temporal bone of interest. Four participants underwent FPCT imaging with the HR mode, and 2 participants, with the FF mode.

Reconstruction Parameters
The MSCT dataset was reformatted with 0.6-mm slices every 0.2 mm using a 512 × 512 matrix and a 65–70 mm FOV. FPCT secondary reconstructions were created with the following parameters: manually generated VOI to include only the electrode array; isotropic voxel size, 0.08 mm; 512 × 512 section matrix; sharp image characteristics.

Multiplanar Images
We used MPR to generate 2 oblique 2D images. The first image was in the plane of the posterior SCC and included the 3 electrode contacts of the linear array implanted in the posterior canal and the tip of braided platinum/iridium wire inserted into the common crus. Section thickness was set to 2 mm to include all electrode contacts on 1 image for both planes. Window width and contrast level were adjusted as needed to optimize the visibility of electrode contacts. A 3D representation of the vestibular lumen and vestibular nerve is added in transparency (E and F) to help visualize the anatomy.

Table 1: Demographic information for participants with vestibular implants

<table>
<thead>
<tr>
<th>Participants</th>
<th>Date Implanted</th>
<th>Date Imaged</th>
<th>Age (yr)</th>
<th>Sex</th>
<th>Imaging Protocol</th>
<th>Implant Side</th>
<th>Reference Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>12 Aug 2016</td>
<td>Sep 2016</td>
<td>62, M</td>
<td>MSCT Left</td>
<td>CC</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>4 Nov 2016</td>
<td>Nov 2016</td>
<td>57, M</td>
<td>MSCT Left</td>
<td>CC</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3 Feb 2017</td>
<td>Feb 2017</td>
<td>63, F</td>
<td>FPCT, HR mode Left</td>
<td>CC</td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>15 Dec 2017</td>
<td>Jan 2018</td>
<td>62, F</td>
<td>FPCT, FF mode Left</td>
<td>CC</td>
<td></td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>24 Aug 2018</td>
<td>Sep 2018</td>
<td>51, F</td>
<td>FPCT, HR mode Right</td>
<td>CC</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>31 Aug 2018</td>
<td>Sep 2018</td>
<td>66, F</td>
<td>FPCT, FF mode Right</td>
<td>CC</td>
<td></td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>14 Jan 2019</td>
<td>Feb 2019</td>
<td>53, F</td>
<td>FPCT, HR mode Left</td>
<td>CC</td>
<td></td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>13 Sep 2019</td>
<td>Oct 2019</td>
<td>55, M</td>
<td>FPCT, HR mode Right</td>
<td>SP</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note: CC indicates common crus of the implanted labyrinth; SP, in a subperiosteal pocket outside the temporal bone; Aug, August; Jan, January; Sep, September; Dec, December; Feb, February; Nov, November. a Age in years at time of implantation.

FIG 2. Method for generating MSCT (A and B) and FPCT (C and D) MPR. Two planes are generated. The first plane is approximately tangential to the thin segments of the superior and horizontal SCCs at their junctions with their ampullae and includes the 6 electrode contacts of the forked array inserted into the superior and horizontal ampullae. The second plane is in the posterior plane of the SCC and includes the 3 electrode contacts of the linear array implanted in the posterior canal and the tip of braided platinum/iridium wire inserted into the common crus. Section thickness was set to 2 mm to include all electrode contacts on 1 image for both planes. Window width and contrast level were adjusted as needed to optimize the visibility of electrode contacts. A 3D representation of the vestibular lumen and vestibular nerve is added in transparency (E and F) to help visualize the anatomy.
Subjects 2, 3, 6 reference electrodes inserted in the common crus (not visible, \( n = 0 \); blurred, \( n = 6 \); clearly visible, \( n = 12 \)). A blooming-type artifact (Fig 2A), in which the electrode array appears larger than its actual size, was identified on the MSCT images both between contacts and at the level of individual electrode contacts. Electrode arrays appeared to occupy most of the ampullae space, making it challenging to identify precisely each electrode contact.

FPCT was able to identify individually all 9 stimulating electrode contacts of the MVI arrays in all 6 participants, as well as all 6 reference electrodes inserted in the common crus (not visible, \( n = 0 \); blurred, \( n = 6 \); clearly visible, \( n = 60 \)). A linear sunburst streak artifact (Fig 2C) was observed in all FPCT images and was noticeable at the level of individual electrode contacts but was reduced between electrode contacts. The use of the HR mode when obtaining FPCT images (\( n = 4 \) participants) produced the clearest images of the electrode array and surrounding labyrinthine structures. After we used the FF mode (\( n = 2 \) participants), all individual electrode contacts could be identified, but electrodes and surrounding osseous structures were less well-resolved compared with the HR mode. MPR reconstructions for all subjects are shown in Fig 4.

**Common Crus Insertion Depth**

The median reference electrode insertion depth angle was 9° (range, −57.5° to 45°). The median reference electrode insertion length was 42 mm (range, 21–66 mm). The reference electrode tip was within the common crus in 3 participants, in the superior canal proximal to the common crus in 1 patient, in the posterior canal in 3 participants, and in a subperiosteal pocket in one. Results are summarized in Table 2.

**DISCUSSION**

Accurately positioning each electrode array near the nerve branch of the SCCs that it is intended to stimulate can maximize the strength and selectivity of the prosthetic stimulation because it reduces the current intensity required to achieve and excite a given proportion of neurons in the targeted nerve branch and also reduces current spread to adjacent neurons in other vestibular nerve branches. Vestibular implant outcomes can vary considerably depending on the strength and selectivity of the electrode-nerve interface, as indicated by variation in the magnitude and direction of reflex eye movements driven by stimuli meant to target each nerve branch individually. Electrode distances to target and nontarget nerve branches are key determinants of the strength and selectivity of stimulation; changing electrode location by ∼200 um can change implant outcomes dramatically. Typically, 1 electrode on an electrode array of a given canal outperforms the others that are 250–500 um away.

Knowing vestibular implant array location, their insertion depth, and distance from target vestibular nerve branches can provide helpful information to choose the best electrodes to activate and define stimulus parameters to use, valuable insights that can drive iterative improvements in electrode array design and surgical technique. For example, electrode contact locations can be used as input to individualized finite element models that, once adequately validated via comparison with real data, can facilitate interpretation of empiric data, generation of testable

**RESULTS**

**Visualization of Individual Electrode Contacts**

Individual electrode contacts were barely discernible in the 2 participants imaged using MSCT. Contacts and osseous structures were detectable but blurred enough so that only 12 of the 18 stimulation electrode contacts could be individually identified (not visible, \( n = 0 \); blurred, \( n = 6 \); clearly visible, \( n = 12 \)). A blooming-type artifact (Fig 2A), in which the electrode array appears larger than its actual size, was identified on the MSCT images both between contacts and at the level of individual electrode contacts. Electrode arrays appeared to occupy most of the ampullae space, making it challenging to identify precisely each electrode contact.

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In the present study, we found that the locations of stimulating electrodes and their relation to vestibular bony structures can be depicted precisely with FPCT. All stimulating electrode arrays were close to their target end organs within the target ampullae; however, they varied with respect to location: adjacent to or far from the bone walls of each ampulla. Electrode contact with vestibular labyrinth walls may influence electrode impedance and the spatial pattern of current density, altering stimulation efficiency and selectivity. We also found that reference electrode location varied significantly from case to case, likely because the surgical technique used (making as small an entry as possible in the posterior canal near the common crus, then sliding the reference electrode in with the intent of it reaching the common crus) does not permit direct intraoperative observation or steering of the electrode tip. In 2 cases, the reference electrode was inserted into the superior SCC instead of going down through the common crus. Intraoperative fluoroscopy or DynaCT may be helpful for guiding or confirming the electrode location.

**FPCT versus MSCT**

FPCT is a relatively new imaging technology that implements flat detectors to create volumetric reconstructions. Several advantages have been seen in angiography and temporal bone imaging for cochlear implantation. FPCT is a rapid imaging technique that obtains a full dataset of temporal bone images in approximately 20 seconds. The most clinically significant advantage of FPCT over MSCT is the ability for small voxel areas to be viewed with high resolution. Due to its higher spatial resolution, FPCT yields equal or higher image quality than MSCT when assessing bony structures of diagnostic interest for radiologists. Potential drawbacks to using FPCT for temporal bone imaging compared with MSCT include lack of widespread availability and poorer resolution of soft tissue with currently available FPCT systems. A significant reduction in artifacts was appreciated on FPCT images over MSCT images. A blooming artifact, with the electrode array appearing larger than its actual size, was identified on the MSCT images. This smooth, concentric artifact was identified both between and at the level of the individual electrode contacts, making it challenging to identify precisely each electrode contact. A beam artifact (linear streak bands) was noticeable on the FPCT images at the level of individual electrode contacts but was significantly reduced between electrode contacts. The position of the electrode contacts was better assessed on the FPCT images mainly because of the decrease in artifacts between them.

Previous authors have reported that an important advantage of FPCT is a reduced radiation dose compared with standard

<table>
<thead>
<tr>
<th>Participant No.</th>
<th>CC Insertion Intended</th>
<th>Insertion Depth Angle</th>
<th>Insertion Length (mm)</th>
<th>Anatomic Location of the Reference Electrode Tip</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Yes</td>
<td>−3°</td>
<td>39</td>
<td>Common crus</td>
</tr>
<tr>
<td>2</td>
<td>Yes</td>
<td>+26°</td>
<td>66</td>
<td>Superior canal</td>
</tr>
<tr>
<td>3</td>
<td>Yes</td>
<td>−30°</td>
<td>42</td>
<td>Posterior canal</td>
</tr>
<tr>
<td>4</td>
<td>Yes</td>
<td>−57°</td>
<td>21</td>
<td>Posterior canal</td>
</tr>
<tr>
<td>5</td>
<td>Yes</td>
<td>+21°</td>
<td>46</td>
<td>Common crus</td>
</tr>
<tr>
<td>6</td>
<td>Yes</td>
<td>−2°</td>
<td>41</td>
<td>Posterior canal</td>
</tr>
<tr>
<td>7</td>
<td>Yes</td>
<td>+45°</td>
<td>51</td>
<td>Common crus</td>
</tr>
<tr>
<td>8</td>
<td>No</td>
<td>NA</td>
<td>NA</td>
<td>Outside temporal bone</td>
</tr>
</tbody>
</table>

Note: NA indicates not applicable.

**FIG 4.** MSCT (A–D) and FPCT (E–P) multiplanar reconstructions for all participants. A and B, Participant 1 MSCT. C and D, Participant 2 MSCT. E and F, Participant 3 FPCT. G and H, Participant 4 FPCT. I and J, Participant 5 FPCT. K and L, Participant 6 FPCT. M and N, Participant 7 FPCT. O and P, Participant 8 FPCT. In every panel, the top of the image is superior and the left edge of the image is anteromedial. Sep indicates September; Feb, February; Oct, October; Nov, November; Jan, January.

hypotheses, and optimization of electrode array designs through simulation. In the present study, we found that the locations of stimulating electrodes and their relation to vestibular bony structures can be depicted precisely with FPCT. All stimulating electrode arrays were close to their target end organs within the target ampullae; however, they varied with respect to location: adjacent to or far from the bone walls of each ampulla. Electrode contact with vestibular labyrinth walls may influence electrode impedance and the spatial pattern of current density, altering stimulation efficiency and
Temporal bone MSCT protocols. Depending on the clinical question, FPCT can reduce the radiation dose even more by imaging only the implanted ear. The collimation available in HR mode can direct the x-ray beam to the ear of interest, minimizing the dose received by the head.

Protocol for Vestibular Implant Imaging

Similar to imaging after cochlear implantation, FPCT imaging of postoperative vestibular implantation is easy to perform, produces high-resolution images, and can depict all individual electrode contacts. Patients should be positioned supine with the head pitched to prevent shadowing/overlap of the stimulator and/or magnets with the inner ear. Our imaging methods use a high-resolution secondary reconstruction algorithm with a manually generated small VOI (voxel size = 0.08 mm), Hounsfield unit kernel type, and sharp image characteristic. A 20-second FPCT acquisition of the head is performed using the following parameters: 109 kV, small focus, 200° rotation angle, and 0.4° per frame angulation step. Before image acquisition, the external components of the vestibular implant system are removed. The patient’s head is pitched forward ~20° by flexing the neck and supporting the head on a firm wedge. The head is then taped in place. This positioning ensures that the Reid plane (which contains the center of the external auditory canals and the cephalic edges of the infraorbital rims, both easily palpable landmarks) is pitched ~20° from Earth-vertical, so that the horizontal SCC plane is approximately perpendicular to the gantry rotation axis and scatter artifacts from the magnets and stimulator electronics will not shadow the inner ear (Fig 5). Acquisition should include the entire implant, overlying scalp (to check scalp thickness over the magnets), inner ears, and at least the maxillary teeth (which are used to get canal orientation relative to a bite block for programming the alignment matrix of the processor).

Finally, multiplanar reconstructions are performed in the axial (horizontal plane of the SCC) and coronal planes, the planes of the superior and posterior canals (which also gives slices through the basal turn of the cochlea), and the plane that contains the superior and horizontal forked electrode array. This last plane should be tangential to the superior and horizontal canals at the junction of their ampullae as described in the Materials and Methods above. When reporting imaging findings of FPCT, we recommend that in addition to commenting on common temporal bone imaging findings, additional comments should be made on the quality of the examination, the number of metallic artifacts, the locations of stimulating electrode contacts relative to the ampullae, the location of the tip of the reference electrode if applicable; and scalp thickness over the implant.

This study has some limitations including a small sample size, lack of a control group, and lack of clinical information regarding correlation between vestibular function recovery and electrode contact positions. These topics are beyond the scope of this article, which aims to present the very first imaging results that are currently under investigation to be addressed in future studies in which imaging plays an important role.

Future development of vestibular imaging may lead to intraoperative DynaCT. DynaCT technology can be applied with acceptable additional time requirements without adding too much complexity to the surgical procedure. Intraoperative data acquisition by DynaCT may represent a suitable option for real-time surgical navigation during a vestibular implant operation. This imaging technology will encourage further advances in vestibular implant surgery and integrate functional aspects of imaging by applying individualized anatomy-based mathematic models that will help predict vestibular flow current spreading for each patient to further understand clinical outcomes of prosthesis implantation.

CONCLUSIONS

FPCT produces high-resolution images of vestibular implants, allowing identification of individual electrode contacts and quantification of their locations relative to vestibular SCC ampullae. Reduced artifacts were seen in FPCT images compared with MSCT images. Optimal FPCT imaging includes a high-resolution secondary reconstruction algorithm with a manually generated VOI that includes only the electrode array. As MVI imaging improves, so will our understanding of the relationships among vestibular anatomy, MVI electrode placement, vestibular performance, and hearing outcomes.
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