INTRODUCTION

Implantable acoustic prostheses can be classified into three categories according to the mechanism of inner ear stimulation. The first category comprises middle ear implants (MEI), which transmit an amplified vibration to the cochlea via the ossicular chain [1-7]. A second category comprises bone-conduction implants that transmit vibration via the bone, excluding the middle and outer ear. The third category comprises prostheses that stimulate the cochlea by vibrating the perilymph through the oval window (OW) for severe-to-profound mixed hearing loss, and are designated as direct acoustic cochlear implants (DACI). The concept was presented initially by Hausler et al. [8] in 2008 as a percutaneous system, called DACS, and was further developed into the transcutaneous Codacs system (Cochlear Ltd., Sydney, Australia), which was investigated in two clinical trials [9,10] and is currently commercially available.

The Codacs system comprises an implantable part, consisting of a receiver coil, the implant electronics, and the electromagnetic actuator, which is held in place by a fixation system, and an externally worn part that is a behind-the-ear sound processor with a radio-frequency coil, i.e., the Cochlear Nucleus CP810 sound processor with modified firmware that facilitates acoustic signal processing. The actuator has a terminal—referred to as the ‘artificial incus’—which is similar to the size of the long process of the incus. The fixation system is fastened to the mastoid cortex with screws, with the ball joint allowing movement and locking of the actuator into the correct position [11]. The artificial incus should be positioned in such a way that it is aligned to the level of the natural incus, above the OW, while avoiding contact with the surrounding tissues and bony structures.

The surgical technique used to insert this system when it was first developed was similar to that used for the minimally inva-
The direct acoustic cochlear implant (DACI) is a innovative and useful implantable prosthesis. We evaluated a new surgical procedure to place the DACI. We proposed to implant the DACI through the superior-anterior tympanotomy, in place of the incus.

**Fig. 1.** The two surgical procedures. (A) Haeusler and Lenarz procedure: device in the mastoid with the artificial incus, through the posterior tympanotomy, orthogonal at the incus. (B) Our procedure: device in the mastoid with the artificial incus, through the anterior-superior tympanotomy, in place of the incus.

The surgical procedure included a small mastoidectomy, limited to a retro-canal tunnel and a posterior-inferior tympanotomy, with access to the facial recess at the level of the OW. This access allowed introduction of the fixation system and the actuator, positioning of the artificial incus, and execution of a stapedotomy. In the operations performed by Hausler et al. [8] in four patients with severe-to-profound hearing loss due to otosclerosis, two stapes prosthesis were used: one anchored to the artificial incus and the other to the natural incus. Hausler et al. [8] reported good functional results; in particular, a gain in the postoperative air conduction threshold with the implant inactive and the piston anchored to the natural incus (range, 14 to 28 dB; n = 4) and a markedly higher gain with the implant active (range, 41 to 62 dB; n = 4). The intervention took a relatively long time (range, 2.5 to 5 hours; n = 4). It was technically complex, but without any noticeable intra- or postoperative complications. The risk of cochlear damage was comparable to that of a classical stapedotomy [8]. In 2012, Zwartenkot et al. [13] compared the results of these four patients to those obtained with nine Vibrant Soundbridge (VSB) implants and nine middle ear transducer (MET) implants. The DACS system demonstrated a better amplification capacity than the VSB and was similar or superior to the MET, but the surgery required was considered more invasive, as it included opening of the vestibule.

More recently, Lenarz et al. [9,10] published the results of two multicenter studies with the DACI system. The surgical procedure was based on the approach used by Hausler et al. [8], but none of the ossicular chains was reconstructed with the second stapes prosthesis. Similar to Hausler et al. [8], Lenarz et al. [9,10] placed the device in the mastoid, with the tip on the footplate of the stapes, through the posterior tympanotomy (Fig. 1). This posterior approach to the middle ear leads to a reduced view of the stapes footplate, the need to expose the facial nerve, and sometimes the need to drill into a part of the posterior canal wall. Moreover, the reduced view of the stapes footplate obligates the surgeon to make a hole in the stapes footplate before placing the device. Therefore, the vestibule stays open for about 30 minutes to 1 hour. The relative complexity of the surgery was highlighted and, in the first trial, the average surgical time was 4 hours and 52 minutes (SD = 1 hour and 26 minutes), including intraoperative testing with a laser Doppler vibrometer.

In consideration of the technical difficulties encountered with the placement of this system, which are reported by all surgeons, we sought to find a simpler implantation procedure. Here, we report the experimentally validated feasibility of a new, simpler, and faster approach for DACI placement.
MATERIALS AND METHODS

We developed a new surgical procedure for implanting the DACI on five frozen temporal bone specimens in the laboratory of the ENT unit of the University Hospital of Pisa, which was equipped with a surgical microscope (Leica M400; Leica, Wetzlar, Germany), as well as surgical drill and instruments. The human anatomical samples consisted of the bone and the soft tissue of the temporal and mastoid region. Ethical approval was waived by the University of Pisa.

Before the intervention, a high-resolution computed tomography (CT) scan of the bones was performed to establish the dimensions of the mastoid and the epitympanic region. The CT scan, with a 0.6-mm section thickness, allowed axial, coronal, and sagittal reconstruction of temporal bone images. All the surgical procedures were performed by a senior surgeon.

Surgical procedure

The dissection begins with the canal incision of a tympanomeatal flap, its detachment, and drilling into a part of the epitympanic frame sufficiently large to allow access to the OW, as performed in stapedioplastic surgery. Through this access, at the end of the surgical procedure, the final surgical steps of removal of the stapes superstructure, stapedotomy, and accommodation of the stapes prosthesis are performed.

The retroauricular access through the soft tissue is similar to that commonly used in the surgery for MEI and DACI. The retroauricular access through the bone starts with a partial mastoidectomy, limited to the superior cell groups. The tegmen, anterior epitympanum, the anterior-superior wall of the external ear canal (EAC), the malleus head, and the body of the incus are identified. The wall of the EAC is thinned, the incus is removed after being disjointed from the stapes, and the bone above the second portion of the channel of the facial nerve is drilled until the OW can be identified (superior-anterior tympanotomy).

The fixation system is inserted and secured to the mastoid with bone screws. The artificial incus is positioned above the OW through the superior-anterior tympanotomy. The ball joint can be adjusted so that the correct position for the artificial incus can be achieved.

The artificial incus has no contact with the malleus, with the edges of the tympanotomy, or with the posterior wall of the tympanic membrane. Its final position is similar to that of the natural incus (Fig. 1). We used the stapes head to find the correct position for the artificial incus. The artificial incus was placed in the same position as the natural incus (Fig. 2A).

The drilling for the receiver coil, implant electronics, and the connecting cable is similar to that used in cochlear implant surgery. The removal of the stapes superstructure, the stapedotomy and the insertion of the stapes prosthesis are performed via the ear canal at the end of intervention (Fig. 2B). Table 1 presents the main steps of the Haeusler and Lenarz procedure and our procedure, whereas in Table 2, we show the main differences between the two procedures [8,9].

RESULTS

We successfully implanted the device in the five anatomical samples, following the new surgical procedure. All surgical steps were found to be fast and simple to execute. The average time for placement of the DACI in the temporal bones following this new procedure, including the dissection of the soft tissues and the suture, was 1 hour and 52 minutes (range, 1 hour and 34 minutes to 2 hour and 22 minutes). Large differences were not
observed among the five temporal bones. In the 3 large mastoids, our procedure was simpler than that in the 2 small temporal bones. The learning curve was crucial, as in fact the last two surgical procedures were the shortest.

**DISCUSSION**

The DACI system is innovative as it transmits the vibrations directly to the labyrinth through a stapes prosthesis, completely bypassing the ossicular chain. The DACI system was designed to be used in patients with severe-to-profound mixed hearing loss. For this indication, the DACI system is significantly applicable, as it constitutes the only alternative to the cochlear implant, and is much less invasive [8-10,13]. The clinical trials conducted with the DACI system to date appear to demonstrate that the risk to the cochlea is comparable to that of a stapedioplasty, while suggesting it may be necessary to investigate the procedure in a larger number of patients, with longer term follow-up, to fully assess the effects of vibration of the prosthesis in the labyrinth [8-10]. Given the relatively recent implementation, DACI placement surgery is still evolving, even with regard to some relevant aspects of the surgical technique. The surgical technique practiced by Lenarz et al. [9] differs from that originally indicated by Hauser et al. [8] in terms of the use of a single stapes prosthesis, a wider mastoidectomy, and accessing the OW/middle ear through a combined transmastoid and transcanal approach in cases where the anatomy of the mastoid does not allow a good view of the OW through the posterior-inferior tympanotomy alone. These changes have simplified the technique used and reported by Hauser et al. [8] somewhat, and have also enhanced its safety. Nevertheless, the adaptations described by Lenarz et al. [9] have highlighted the difficulties and the long time required for this surgical intervention.

The modifications to the surgical technique proposed in this paper are designed to contribute further to the simplification of the surgical technique and to the reduction of clinical risk and surgical time by decreasing the risk of facial damage, both in normal anatomy and in patients with facial nerve or inner ear anomalies. The main aspects of the changes are the combined transmastoid and transcanal access to the OW, the partial mastoidectomy, anterior-superior tympanotomy, the removal of the incus, and the stapedotomy with a single stapes prosthesis anchored on the artificial incus. The combined access to the OW allows performance of the stapedotomy through the ear canal, with the best view of the OW area. Bone drilling is limited to a portion of the posterior-superior tympanic frame as in classical stapedial plastic surgery. This, due to its small size, does not involve reconstruction with cartilage, as in operations with combined access performed by Lenarz et al. [9]. Mastoid drilling is limited to the superior cell groups and to the anterior epitympanum. In our experiment on the temporal bone, the removal of

| Table 1. Description of the main steps of the Haeusler and Lenarz procedure and our procedure |
|-----------------------------------------------|
| **Haeusler and Lenarz procedure** | **Our procedure** |
| Retroauricular skin incision | Retroauricular skin incision |
| Mastoidectomy | Little mastoidectomy in epitimpanic area |
| Enlarge posterior tympanotomy | Transcanal access to visualize the stapes |
| If the stapes is not well visualized, drill a part of the posterior canal wall or transcanal access | Small anterior-superior tympanotomy |
| Remove the crura of the stapes and generate a hole in the footplate | Remove the incus |
| Prepare the bone bed and place the fixation system | Prepare the bone bed and place the fixation system |
| Prepare the bone bed and place the processor | Prepare the bone bed and place the processor |
| Place the device in the mastoid with the artificial incus through the posterior tympanotomy, orthogonal at the incus | Place the device in the mastoid with the artificial incus through the anterior-superior tympanotomy, in place of the incus |
| Place the stapes piston (through the posterior tympanotomy) | Remove the crura, make the hole in the footplate and place the stapes piston (through the canal) |

The main difference between the two procedures are shown in bold.

| Table 2. Main differences between the two procedures |
|-----------------------------------------------|
| **Main differences** | **Haeusler and Lenarz procedure** | **Our procedure** |
| Time spent for surgery | Approximately 4 hr | Approximately 2 hr |
| Risk of facial and corda tympani damage | Low (risk of a large posterior tympanotomy) | Very low (posterior tympanotomy is not performed) |
| Risk of cochlear damage | High (the vestibule stays open for approximately 30 min to 1 hr, the hole in the footplate is made before placing the device) | Very low (the vestibule stays open for a few seconds or minutes, the hole in the footplate is made during a normal stapedioplasty) |
| Incus removal | No or only occasionally | Always |
the incus and the anterior-superior tympanotomy allowed easy access to the OW for the introduction of the actuator, under good control for the positioning of the artificial incus. This tranmastoid access, which ends with a anterior-superior tympanotomy, differs from those practiced by Hauser et al. [8] and by Lenarz et al. [9,10], which require a rather large posterior-inferior tympanotomy. The type of access proposed in this paper, however, involves the removal of the incus.

Hauser et al. [8] used the incus for the placement of a second stapes prosthesis in the OW, in addition to the stapes prosthesis anchored on the artificial incus. The functional results show that the contribution of the second stapes prosthesis is insignificant compared to that of the stapes prosthesis placed on the artificial incus. None of the later surgeries practiced by Lenarz et al. [9,10] used the second stapes prosthesis in the OW, thereby reducing the cochlear risk as a consequence of the stapedectomy. On the other hand, the presence of both the natural incus and artificial incus occupies space near the OW, complicating the positioning and the stapedectomy. The presence of the natural incus, therefore, does not seem to play a favorable role in DACI placement and/or performance outcomes. Even in the event of a DACI failure, and a decision to opt for a CI surgery, it would be hard to conceive of a possible role for the natural incus. As stated in the instructions for use, the partial removal of the incus may be required as part of the surgical technique for implanting the DACI.

In conclusion, the purpose of our study was to examine the feasibility of an amended DACI placement procedure to both facilitate and expedite the standard procedure currently used. We tested this procedure on a small number of samples (five temporal bones), similar to the number of samples originally reported by Hauser et al. [8] for the minimally invasive cochlear implant technique. Our proposal for an amended DACI surgical procedure takes into consideration the combined access to the OW, transtympanic for the stapedotomy, and transcervical access with the removal of the incus, and an anterior-superior tympanotomy for accommodation of the actuator and the artificial incus. This simplified the surgical procedure and consequently reduced the time required for DACI placement, and it reduced the risk of facial and corda tympani damage; however, the actual effects should be verified in patients.

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CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.