Purpose: Electronic retinal implants restore some visual perception in patients blind from retinitis pigmentosa. Eye movements cause mechanical stress in intraorbital power supply cables leading to cable breaks. By using computer tomography (CT) scans at the extreme positions of the four cardinal gaze directions, this study determined in vivo, which of three surgical routing techniques results in minimal bending radius variation and favors durability.

Methods: Nine patients received the first-generation subretinal implant Alpha IMS (Retina Implant AG, Reutlingen, Germany) in one eye. Three techniques for intraorbital cable routing were used (straight cable route (A), parabulbar loop (B), and encircling band (C)), each in three patients. All patients underwent computer tomography of the orbital region. The bending radius of the intraorbital cable was measured with the DICOM viewer Osirix v4.1.2 (Pixmeo SARL, Bernex, Switzerland) and served as indicator for mechanical stress.

Results: Average bending radius variation was 87% for method A, 11% for method B, and 16% for method C. Methods A and B ($P = 0.001$) and methods A and C ($P = 0.001$) differed significantly, while method B and C showed no statistical difference ($P = 0.074$).

Conclusions: Compared to straight routes, arcuated cable routes significantly reduce cable movement and bending. Due to an easier surgical procedure, a parabulbar loop is the preferred method to minimize bending radius variation and prolong survival time of electronic subretinal implants.

Translational Relevance: CT analysis of cable bending of implanted medical devices allows to determine which surgical routing technique favors durability in vivo.

Introduction

One in 4000 of the world population suffers from retinitis pigmentosa (RP), a major cause of visual disability and blindness in middle-aged people. Initially, the neuronal degeneration is limited to the first neuron of the visual system (the photoreceptor), whereas many of the second neurons (bipolar cells) and third neurons (ganglion cells) still operate for a long time during the course of disease. One way to partially restore visual function is the use of a...
subretinal prosthesis Alpha IMS (Retina Implant AG, Reutlingen Germany), which replaces the function of photoreceptors (Fig. 1).3–5

The implant is driven by a battery and electronic control elements in a handheld unit, with a wireless transdermal connection by a retroauricular coil system.3,6 This system provides power and enables the patient to adjust sensitivity and contrast settings of the system via electromagnetic induction.5

Given two to three saccadic eye movements per second with additional microscopic, fixational eye movements in between, the intraorbital parts of the implant's power supply cables, which connect the subretinal implant to the retroauricular coil, are exposed to considerable mechanical stress.7,8 As patients blind from RP are comparably young and have a long lifespan ahead, durability and suitability of implant parts are essential.1 Hafed et al. demonstrated in postoperative eye tracking experiments in patients with Retina Implant Alpha, that eye movements for gaze can recover.9

Therefore, minimizing cable movement and, simultaneously, mechanical stress to the cable and the surrounding tissue is crucial, especially as cable breaks during initial phase of trials with Alpha IMS were common.3 Such cable problems were solved in the subsequent model, the Retina Implant Alpha AMS,10 after extensive preclinical testing, outlined in the discussion section.

There are various implants that avoid orbital cables, for example, the PRIMA device11; instead of intraocular electronic amplifiers relatively bulky electronic goggles for amplification of image intensity need to be employed. Such goggles, however, reduce the utilization of natural gaze for object detection; not the case for patients after having received Retina Implant Alpha devices.9

The present study focuses on patients with the Retina Implant Alpha IMS and compares the variation in bending radius of the intraorbital cable during eye movements in three different routing techniques (straight cable route (A), parabulbar loop (B), and encircling band (C)) in the extreme positions.

**Materials and Methods**

**Subjects**

Nine patients (four females, five males), mean age ± standard deviation (SD) = 46.9 ± 7.2 years (age range 35–62 years) received the subretinal implant Alpha IMS (Retina Implant, Reutlingen, Germany) in the first single center part (2010–2011) of a multicenter trial (www.clinicaltrials.gov, NCT01024803). Prior to surgery, visual function was severely reduced due to legal blindness (eight patients: light perception without correct light source localization; one patient: no light perception) due to either RP (eight patients) or cone-rod dystrophy (one patient). None of the patients had additional eye disorders affecting the ascending optic pathways. Prior to study participation, all participants gave written informed consent in conformity with the Declaration of Helsinki. The local ethics committee approved the study, and it was performed in accordance with the German Medicinal Product Law (MPG) and EN ISO 14155.12
Figure 2. Computer tomographic reconstruction of the retroauricular position of the ceramic box, containing the coil and the electronics that enable to transmit energy and to control the implant’s stimulation parameters, and the silicon cable running from the box to the orbit (Subject RIAG TU07).

Electronic Retinal Implant (“Retina Chip”)

An integral part of the electronic retinal implant is a complementary metal–oxide–semiconductor (CMOS) chip, which has approximately 1500 pixels. Each pixel contains a photosensitive diode, an amplifier, and a stimulation electrode. The CMOS chip is mounted and connected to a flexible polyimide foil (intraocular part) that is connected to a silicon cable (extraocular part) via a ceramic adapter plate fixated on the sclera (Fig. 1). The silicon cable connects the chip via the adapter plate to the power supply in a special ceramic box compartment, which is further connected to a subdermal reference electrode behind the ear. This ceramic box compartment contains various electronic parts and a magnetic coil receiver for wireless power transfer through the skin (Figs. 1 and 2). This system is implanted completely subdermally. The power from an external battery compartment is transmitted wirelessly through the skin using two inductive/magnetic coils, one in the ceramic box compartment and one external coil kept in place by a magnet.

Implantation/Surgical Procedure

The electronic subretinal implant was implanted as described previously. In brief, the tip of the polyimide foil containing the CMOS chip with the actual photosensitive pixels was implanted into the subretinal space, preferably under the fovea at the posterior pole of the eye (Fig. 1). The polyimide foil with the chip on its tip was inserted into the subretinal space toward the fovea from a superior lateral scleral incision near the equator of the eye (Fig. 1). The ceramic adapter plate that uses six gold wires to attach the polyimide foil to the round silicon cable was sutured to the sclera. From this first fixation point on the moving eyeball, the silicon cable was running to another fixation point at the orbital rim of the upper temporal orbit (Fig. 3). The silicon cable then runs subperiostally from the orbital rim to the retroauricularly implanted ceramic box compartment (Fig. 2).

Between the two fixation points in the orbit, the silicon cable must follow the eye movements, while providing a stable electrical connection between the
intraocular implant and the power supply system in the retroauricular ceramic box compartment. This is accomplished using one of the following three different routing techniques (Fig. 4)\textsuperscript{12}:

**Method A:** The silicon cable runs straight on a direct short route from the orbital rim to the eyeball with only enough cable length to allow for limited eye movement, mainly in central viewing direction (maximum elongation of the silicon cable, Fig. 4A).

**Method B:** The silicon cable forms a parabulbar loop, thus distributing the movement onto a longer cable segment (Fig. 4B).

**Method C:** The silicon cable forms a loop around the eyeball (similar to scleral buckle surgery, which is used for retinal detachments) to reduce transmission of mechanical forces onto the intraocular parts of the implant (Fig. 4C).

**CT Imaging Technique**

Video fluoroscopy (7.5 p/s; 55 nGy/p) was performed with a biplane angiography unit Axiom Artis zee (Siemens Healthcare, Erlangen, Germany) in three of nine patients (one patient per surgical method A, B, and C, respectively; 55–65 mGy per patient) to verify the assumption that the majority of the cable movement and therefore bending of the cable happened between the orbital rim and the eyeball, and that the other parts/sections of the silicon cable remain static during eye movement (Videos S1–S6 in supplementary material)\textsuperscript{12,18}.

All nine patients underwent computer tomography examinations (16 × 0.75, 130 mAs, Pitch 0.55; 120 kV) with a Somatom Sensation 16 multislice computer tomograph (Siemens Healthcare, Erlangen, Germany). An individual CT scan of the orbital region was performed for each of four viewing directions, a total of four scans per patient. CTDI (computed tomography dose index) was <25 mGy per scan, totaling <100 mGy per patient. No contrast agent was used\textsuperscript{12,18}.

The four gaze directions were the following: upper temporal, upper nasal, lower temporal, and lower nasal gaze direction (Fig. 5). Diagonal gaze directions were chosen to exploit maximum extension as well as maximum bending radius of the cable running diagonally from the upper temporal orbital rim to the eyeball\textsuperscript{12,18}.

**Determination of Cable Bending Radius**

Datasets of each scan were analyzed in the 3D multiplanar reconstruction mode with the free DICOM viewer Osirix v4.1.2 (Pixmeo SARL, Bernex, Switzerland). One viewing plane was aligned precisely to the plane of the silicon cable route to avoid projection errors during measurement. The Circle ROI Tool (Oval ROI Tool while holding shift key) was aligned to the apex of the cable curve, showing the area of the circle ROI (Figs. 6 and 7). The radius of the circle outline was calculated via the formula:\textsuperscript{12,18}

\[
radius = \sqrt{\frac{\text{area}}{\pi}}
\]

Mean and standard deviation (SD) of the bending radius was calculated for each subject. In particular, the determination of SD of the bending radius is important to estimate the mechanical stress on the cable: Minimal change of bending radius during eye movements causes minimal stress, whereas huge changes cause maximal stress, regardless of the mean bending radius in a moving eye\textsuperscript{12,18}.

We performed repeated measurement analysis of variance (rmANOVA) to compare the three different routing techniques with the within-subject factor viewing direction (four levels: upper temporal, upper nasal, lower temporal, and lower nasal) and the
Figure 5. Cable position during eye movement for (A) straight route (method A, Subject RIAG TU05) and (B) encircling band (method C, Subject RIAG TU07), (C) parabulbar loop (method B, Subject RIAG TU12) four diagonal viewing directions each: (a) nasal superior (b) nasal inferior (c) temporal superior, and (d) temporal inferior.
between-subject factor operation method (three levels: method A (straight), method B (parabulbar loop), and method C (encircling band)). In case of significance, we used paired and two-sample t-test for further analysis.12,18

**Operating Times**

We compared the operating times for the three different surgical methods using analysis of variance (ANOVA) with the inner-subject factor operation method.
Table 1. Individual Bending Radius

| Bending Radius (mm) | Method A (Straight Route) | Method B (Parabulbar Loop) | Method C (Encircling Band) |
|---------------------|---------------------------|----------------------------|---------------------------|
|                     | Sub 2 Sub 6 Sub 5         | Sub 9 Sub 12 Sub 1         | Sub 7 Sub 8 Sub 10        |
| Upper temporal (mm) | 2.3 0.6 1.4               | 2.7 1.9 2.5                | 5.1 3.9 5.5               |
| Upper nasal (mm)    | 3 9 6.8                   | 2.7 1.8 2.2                | 4 3.7 4.2                 |
| Lower temporal (mm) | 3.1 2.1 5.9               | 3.1 2.3 2.2                | 4.7 3.1 3.9               |
| Lower nasal (mm)    | 16.6 9.2 11               | 3.1 2.4 2.7                | 3.6 4.4 3.6               |
| Mean ± SD (mm)      | 6.3 ± 6.9 5.2 ± 4.5 6.3 ± 3.9 | 2.9 ± 0.2 2.1 ± 0.3 2.4 ± 0.2 | 4.4 ± 0.7 3.8 ± 0.5 4.3 ± 0.8 |
| Standard Deviation (%) | 111 86 63               | 8 14 10                    | 16 14 19                  |
| Mean variation of bending radius (%) | 86.7 ± 24.0          | 10.7 ± 3.1                | 16.3 ± 2.5                |

Table displays the individual bending radius in mm for nine subjects in three different surgical techniques (A, B, and C) in four viewing directions (upper temporal, upper nasal, lower temporal, lower nasal) and the mean bending radius for each subject as well as the standard deviation. The variation of bending radius for each method is calculated. Percentages indicate the standard deviation relative to the mean for each subject. The change in bending radius for each method is calculated from the mean of percent standard deviation for the three participants, who were operated with the corresponding method.

Statistical Analysis

IBM SPSS Statistics 25.0 (International Business Machines Corporation, Armonk, NY, USA) was used. Whenever the Shapiro-Wilk test indicated a deviation from normal, we used logharitimized data. Whenever Mauchly’s Test of Sphericity was violated, we used the Greenhouse-Geisser correction. A $P$-value of $< 0.05$ was regarded as statistically significant. The Bonferroni correction was used to correct for multiple comparison ($P < \frac{0.05}{3} = 0.017$).

Results

Optimal Bending Radius

Average bending radius ± SD was 5.9 ± 4.8 mm for method A ($n = 3$ patients), 2.5 ± 0.4 mm for method B...
Intraorbital Cable Movement

Table 2. Operating Times

| Method A (Straight Route) | Method B (Parabulbar Loop) | Method C (Encircling Band) |
|--------------------------|-----------------------------|----------------------------|
| Individual operating time|                            |                            |
| sub 2                    | 480 min                     | sub 1                      | 505 min |
| sub 5                    | 485 min                     | sub 9                      | 595 min |
| sub 6                    | 480 min                     | sub 12                     | 480 min |
| Mean                     | 481.67 min                  | 493.33 min                 | 483.33 min |
| Standard deviation       | 2.89 min                    | 12.58 min                  | 22.55 min |

Table displays the individual and mean operating times for nine subjects in the three different surgical techniques: method A (straight route), method B (parabulbar loop), and method C (encircling band).

Discussion

This study applied computer tomography to determine the cable routing technique for subretinal implants that provides minimal bending of extraocular cables and, hence, might produce minimal mechanical stress due to eye movements in vivo. Reduction of mechanical stress is crucial for long-term function and reliability for daily use of these devices.

We compared the amount of bending radius variation in three different surgical techniques. Method B (parabulbar loop) and method C (encircling band) showed comparable changes in bending radius and might consequently cause similar mechanical cable stress during eye movements (Fig. 8). Both methods showed significantly less cable movement than method A (straight route, Table 1, Fig. 8). Overall, our study demonstrated that both a parabulbar loop (method B) and the more complex encircling band (method C) showed minimal variation in bending radius and might reduce cable stress in subretinal electronic implants due to eye movements.12 Mean operation time was comparable for all methods (Table 2). The whole procedure (intra- and extraorbital surgery) takes about eight hours. However, interindividual variability of surgery time was bigger for methods B and C compared to method A (Table 2), which reflects the higher complexity of the surgical procedures.

The presented method for analysis of implanted cables is not only applicable to subretinal implants, but to any cable structure, exposed to movement within the human body, for example, cerebral shunts used for hydrocephaly treatment or limb prostheses.19–22 Assessment only requires standard CT devices and any DICOM viewer capable of freely adjustable viewing planes and basic measurement tools.

Compared to other organs, the high mobility and permanent intentional and unintentional position changes of the eye exposes cables to severe mechanical stress and can cause changes in microchip position.7,23 In performing CT-imaging, this study revealed that the variation of bending radius between different eye movements can be considerably minimized by optimizing bended cable routings. Kuehlewein et al. had analyzed the change of chip position in patients with subretinal implant Alpha IMS and AMS, including the cohort of this paper, in vivo. All our patients with
straight cable route and parabulbar loop had stable or minor variations of chip position. In contrast, two patients with encircling band had significant changes in chip position and in the third patient the retinal implant had to be repositioned after 48 days due to a retinal hole at the distant border of the chip.23

Cable bending and the optimal course for the orbital cable portion had been investigated in human cadaver head studies with mock surgeries in the anatomy department (K.U. Bartz-Schmidt and F. Gekeler, Tübingen, personal communication) before any clinical application. However, the individual orbital situation of patients in the first clinical trial with the Retina Implant Alpha IMS had required personalized adaptation of the initial cable procedure during the trial and resulted in the various loop sizes described here. Subsequently, Daschner et al. had recorded clinical reliability data in a laboratory set up for advanced aging experiments.10 The power supply cable was tested using machine that simulated the movements of the eye by bending the fixed cable over 27 million times. CT images had been used to optimize the apparatus,24 rendering application of cable bound implants safe.

Indeed, real-life data confirmed that the use of the surgical technique of a parabulbar loop (method B) and material improvements of the cable could eliminate cable breaks in the subretinal implant Alpha IMS and its successor Alpha AMS.3,4 Furthermore, in vivo data revealed that this surgical technique only leads to minor changes in chip position.23 Additional technical improvements in design and manufacturing of the intraorbital cable used in Alpha AMS even prolonged the mean lifetime of the subretinal implant to seven years (compared to 1.5 years in Alpha IMS).10

Consequently, bended cable routings can reduce the mechanical stress on the cable and surrounding tissue and might prolong the implant’s lifetime. This finding is important as subretinal implant patients generally are young and have a long life ahead.1,3,5

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**Supplementary Material**

Videos for method A (Video S1 & S2) and method B (Video S3 & S4).

Supplementary Video S1. Straight route viewed from the bottom left (Subject TU_05).

Supplementary Video S2. Straight route viewed from the front (Subject TU_05).

Supplementary Video S3. Encircling band viewed from the bottom left (Subject TU_07).

Supplementary Video S4. Encircling band viewed from the front (Subject TU_07).

Supplementary Video S5. Parabulbar loop viewed from the bottom left (Subject TU_12).

Supplementary Video S6. Parabulbar loop viewed from the front (Subject TU_12).

Amended July 27, 2021: When the article was first published, there were minor typographical errors in the Abstract. In the Results section, the sentence “Methods A and B (P = 0.005) and methods A and C (P = 0.007) differed significantly, while method B and C showed no statistical difference (P = 0.07)” has now been changed to “Methods A and B (P = 0.001) and methods A and C (P = 0.001) differed significantly, while method B and C showed no statistical difference (P = 0.074).”