Temporal Properties of Visual Perception on Electrical Stimulation of the Retina

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PURPOSE. To investigate the elementary temporal properties of electrically evoked percepts in blind patients chronically implanted with an epiretinal prosthesis.

METHODS. Nine subjects were presented with isolated stimuli of variable duration and pulse rate. Stimulation amplitude was set to the upper comfortable level and a group of 2 × 2 adjacent electrodes was simultaneously activated. First, subjects were asked to verbally describe their visual perception paying particular attention to the time-course of brightness. Then, in subsequent trials, they described the brightness time dependence using a joystick while auditory feedback of joystick position was provided.

RESULTS. All subjects described a bright, well-localized percept at stimulus onset. Only one subject reported such a bright, well-localized visual sensation during an entire 10-second stimulation trial. For the remaining eight subjects, it faded more or less rapidly (in four cases <0.5 second) and was often followed by a percept described as less bright, poorly localized, and having different color. Only initial percepts at stimulation onset seemed bright and localized enough to reconstruct a patterned image. Changing stimulation pulse rate influenced the time course of perception only in some cases but the effect was not systematic.

CONCLUSIONS. Percepts differed considerably across subjects, probably because of the considerable variations in the progression and remodeling processes associated with the disease. Appropriate coding of a patterned image under such conditions appears challenging. Further research of the underlying mechanisms of visual perception upon electrical stimulation of the retina is required to optimize stimulation paradigms and to better establish patient selection criteria.

(ClinicalTrials.gov number, NCT00407602) (Invest Ophthalmol Vis Sci. 2012;53:2720–2731) DOI:10.1167/iovs.11-9344

The first efforts to develop an electronic visual prosthesis started in the late 1960s.1–4 Since then, different approaches for restoring vision via electrical stimulation have been proposed. Among these, retinal prostheses are probably the most advanced approach, as demonstrated by ongoing human clinical trials.

Electrical stimulation of the retina is envisioned as a promising means for restoring some kind of visual perception to blind patients suffering from degenerative diseases of the retina, such as retinitis pigmentosa (RP) and age-related macular degeneration (ARM).5,6 In these diseases, the light-sensitive cells in the retina (photoreceptors) are lost while second-order retinal neurons (bipolar and ganglion cells) are relatively preserved.7–10 Thus, an electrode array implanted on the inner (epiretinal implant) or outer (subretinal implant) retinal surface could be used to directly stimulate the surviving cells and attempt to transmit an “artificial image” to the brain.

Significant research efforts have paved the way from the initial concept to the development of prototypes ready to be tested in human clinical trials (see e.g., references 6, 11–16). The feasibility of the approach was established through acute in vivo experiments on normally sighted subjects and blind patients. The first studies yielded encouraging results.17–19 Electrical stimulation was delivered to the surface of the retina under local anesthesia and visual percepts were successfully elicited in all patients tested. In general, the localization of percepts corresponded well to the site of stimulation and the relationship between the pattern of electrical stimulation and the perception induced.20,21 Despite important intersubject variations, this study yielded similar basic proof-of-concept results. These studies were followed by substantial technical efforts to develop devices adequate for chronic human use.

To date, five groups have launched human chronic clinical trials: (1) Optobionics, Inc.22–25 (Palo Alto, CA) carried out the first attempts of implantation on human volunteers. Improvement of visual perception and/or slowing of vision loss were reported in areas adjacent and distant to the implant. Only 4 of the 10 implanted patients reported intermittent “phosphenelike lights” at the actual location of the implant. These results combined with animal studies24 suggested that this device induced some kind of neurotrophic effect, but that the improvements in visual function observed were unrelated to electrically evoked visual percepts. (2) Retina Implant AG25 (Reutlingen, Germany) led a clinical trial during which 11 blind patients were implanted with a subretinal prosthesis for a period of 4 months. The device consisted of an array of 1500 microphotodiodes (each with its stimulation electronics) and another array of 16 externally controlled (wired) electrodes allowing for direct stimulation of the retina. Results of psychophysical testing have been reported for three patients. All three were able to perform simple visual tasks, such as discriminating the orientation of a group of four adjacent electrodes stimulated simultaneously (e.g., horizontal, vertical,

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Supported by the Swiss National Fund for Scientific Research (Grant 3152001114152) and by the “Suzanne” Funds.

Submitted for publication December 17, 2011; revised March 1, 2012; accepted March 12, 2012.

Disclosure: A. Pérez Fornos, None; J. Sommerhalder, None; L. da Cruz, None; J.A. Sabel, None; S. Moband-Said, None; F. Hafezi, None; M. Pelizzone, None

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oblique), detecting light projected onto the microphotodiode array, and localizing bright large objects (e.g., dishes) on a dark table. One patient achieved more complex tasks, like identifying large (5–8-cm) single letters and putting them together to form words. (3) IMI Intelligent Medical Implants, GmbH (Bonn, Germany; Richard G, et al. IOVS 2008; 49: ARVO E-Abstract 1786) launched another clinical trial designed to test their IRIS system over a 4-month period. This is an epiretinal device containing 49 electrodes and incorporating a “learning” retina encoder that matches the stimulation patterns to those seen by the patient. Unfortunately, little information is available on this trial. Rare public reports (Keserue M, et al. IOVS 2008; 49: ARVO E-Abstract 1785) indicate that no damage to the retina has been observed in implanted patients and that visual percepts have been elicited at charge densities below 1 mC/cm². (4) EpiRet GmbH (Giessen, Germany) conducted a clinical trial designed to evaluate the EPIRET3 visual prosthesis prototype. This epiretinal 25-electrode system was completely implanted within the eye and was tested on six volunteers over a 4-week trial. Safety data and surgical techniques have been presented. Four patients consistently reported visual sensations at stimulation currents below safety limits. When presented with the same stimulation parameters, the description of percepts varied substantially across subjects and three of them were able to achieve simple pattern discrimination tasks. (5) Finally, the largest clinical trial is led by Second Sight® Medical Products, Inc. (Sylmar, CA, Humayun MS, et al. IOVS 2010; 51: ARVO E-Abstract 2022). It is a long-term study (3 to 5 years) offering the possibility of conducting detailed psychological testing on human subjects with electrodes implanted chronically on the retina. The device evaluated is the Argus II epiretinal prosthesis, a second-generation device with 60 retinal electrodes. (The first-generation epiretinal implant by Second Sight® Medical Products, Inc. was the Argus I implant, a 16-electrode device tested on six RP patients. Patients reported discrete phosphenes perception on stimulation and three of them performed better-than-chance on simple visual tasks. The system includes a camera that captures the visual scene and a microprocessor that wirelessly powers an implanted device and controls the currents that are to be delivered to the retina. To date, 32 patients have been implanted worldwide. All patients reported the perception of visual phosphenes on electrical stimulation. Performance results for simple visual tasks, such as localizing a white square presented at random locations on a dark screen and more complex tasks, such as character and word recognition (da Cruz L, et al. IOVS 2010; 51: ARVO E-Abstract 2023; Stanga PE, et al. IOVS 2010; 51: ARVO E-Abstract 426) have been presented. Three “star patients” in the trial have even been able to read short four-word sentences, two of them reaching maximum rates of two to five words per minute (Sahel JA, et al. IOVS 2011; 52: ARVO E-Abstract 3420).

What are the elementary characteristics of visual percepts elicited on continuous electrical stimulation of the retina? This key issue is interesting for our fundamental understanding of the visual system as well as of practical importance for the development of efficient visual prostheses. There is little background information on this, mainly because most of the human studies cited above were of short duration, which limited the amount of data that could be collected. Because our center in Geneva participates in the Argus II clinical trial, we took advantage of the possibility of long-term access to human experimental subjects to study in detail the temporal properties of the visual perception evoked by electrical stimulation of the retina and the influence of some basic stimulation parameters.

**METHODS**

The Argus II Retinal Stimulation System (Second Sight Medical Products, Inc.) comprises both implanted and external elements. The implanted device consists of a 6 × 10 electrode array (200 µm electrode diameter, 575 µm center-to-center spacing) tacked to the epiretinal surface and of a titanium case (attached to the outside of the eye with a scleral band) containing a receiver coil and a microproces-
sor driven stimulator. External components include a body-worn video
processing unit (VPU) and a pair of glasses on which a miniature
camera and a transmitter coil are mounted. Briefly, the image captured
by the camera is processed by the VPU and transformed into a custom
pattern of electrical stimulation. The transmitter coil powers up and
sends commands to the implanted stimulator that finally activates the
retinal electrodes.

The Argus II Retinal Stimulation System Feasibility Protocol
(www.ClinicalTrials.gov NCT00407602) was designed and conducted
in accordance with the Declaration of Helsinki, ICH Guidelines for
Good Clinical Practices (GCPs), ISO 14,155-1:2003, and applicable
local and federal regulations pertaining to medical device clinical trials.
Local approval from the Governmental Health Agencies and from the
Ethics Committee was obtained in each of the countries and
institutions where the study is being conducted. All implanted subjects
had a confirmed history of RP with remaining visual acuity of 2.9
logMAR (measured by an adaptive four alternative forced choice [4FAC]
square wave grating test) or worse in both eyes. Written consent was
obtained from all subjects and the device was implanted in the
patients’ worse-seeing eye. More details on the trial and the Argus II
device can be found in previous publications.

Subject Selection
Nine subjects, selected based on their availability for testing, were
recruited from three European sites participating in the trial: the
Geneva University Hospitals (Geneva, Switzerland), the Moorfields Eye
Hospital (London, UK), and the Quinze-Vingts National Eye Hospital
(Paris, France). Details on the subjects are presented in Table 1.

Experimental Procedure
Subjects were presented with single stimulation trials separated by
long pauses of at least 60 seconds. Single trials consisted in biphasic
pulse trains (cathodic first, 0.46 ms per phase) of variable pulse rate (5,
20, 60 pulses per second [pps]). To complete the characterization of
the time-course of brightness perception, three stimulus durations
were evaluated (1, 10, 60 seconds). A group of 2 × 2 adjacent
electrodes (QUAD) was simultaneously activated and stimulation
amplitude was set to the upper comfortable level (UCL). We used
QUADs instead of single electrodes because they elicited larger visual
percepts, easier for the subjects to describe accurately, and because
their thresholds were lower. For each subject, the tested QUAD was
selected: (1) to have low threshold (i.e., to maximize the available
dynamic range [Please note that the upper safety limit for the system
during psychophysical testing in the clinic] is 1 mC/cm². We never
exceeded this limit in any of the experiments mentioned in the
article.) and (2) to be as close to the fovea as possible. The distance
from the center of the tested QUAD to the fovea is presented in Table 1.

During the initial trials in each experimental condition, subjects
were asked to verbally describe their visual perception paying
particular attention to the time course of brightness. The same
stimulus was repeated as many times as necessary, until subjects felt
comfortable with the words they used for their description. They were
also asked several questions regarding the time course of brightness.

In subsequent trials, subjects were requested to mimic or “plot”
the time course of brightness using a joystick (vertical axis only; see
Fig. 1). The resting (central) position of the joystick corresponded to
“background brightness” perceived in absence of stimulation. The
uppermost (“full push”) position of the joystick corresponded to the
highest brightness level perceived during the whole trial. Positions
below the central position (“pull positions”) were offered to describe
“darker than background” percepts. Joystick position was sampled at
20Hz and mapped to a 0–10 scale, where 10 corresponded to the
uppermost position (highest brightness perceived during the trial) and
0 to “background brightness.” In addition, auditory feedback of
joystick position was provided via a sound of variable pitch (highest
joystick position 3200 Hz, central joystick position 800 Hz, lowest
joystick position 200 Hz).
For each stimulus condition, subjects were allowed to practice ad libitum. Figure 2 presents examples of data collected during the last five trials of a 20-pps, 10-second duration stimulus for S3. The subject systematically perceived a very bright phosphene (10/10 rating) at stimulus onset, but this bright percept lasted only a fraction of the entire stimulus duration. Then, brightness dropped rapidly to 5/10 to 7/10 ratings and slowly faded to background brightness. Stimulus offset was not accurately perceived. As it can be seen from the plots in Figure 2, trial-to-trial reproducibility was remarkable despite the relative complexity of the task. We therefore decided to merge the five last trials collected in each condition and to present averaged data (±SD) in all subsequent results presented in this article.

Finally, to verify the accuracy of subjects in providing a quantitatively precise estimation of brightness with the joystick, they were also asked to provide verbal estimates of brightness in a 1-10 scale at critical time points of the response. Figure 2 shows an example of these brightness estimations for S3, superimposed to the averaged joystick plot (green dots in the bottom right plot). As it can be seen in the graph, this particular subject was quite accurate in matching verbal estimations with joystick data.

**RESULTS**

Figure 3 presents the averaged joystick plots (±SD) of each subject for a 10-second stimulus at 20 pps. They all reported that a well-localized spot in their visual field lit up immediately at stimulus onset. All subjects attributed a brightness level of 10 to this event. However, of the nine subjects tested, only S6 described that this initial well-localized percept remained stable and lasted for the entire duration of the stimulus. For subjects S3, S4, S5, and S8 this initial percept lasted only 2 to 5 seconds, whereas the remaining subjects (S1, S2, S7, S9) experienced a short-duration, flash-like initial percept that lasted less than 0.5 second. Afterward, this well-localized percept “exploded” into a much less localized and lower brightness visual sensation. In addition, some subjects reported a brightness reincrease at stimulus offset that was most often brief (S1, S2, S4) but could also last several seconds (S7). Finally, note that subject S2 described a percept that became “darker than background” upon ongoing stimulation.

The considerable differences observed across subjects cannot be explained by experimental error. First, trial-to-trial reproducibility was very good in all cases (look at the small experimental SDs in each subject’s plot). Second, for every subject, we replicated the same measurements in the same experimental condition in sessions that were several weeks apart. The result was always virtually the same (within experimental error). Finally, we also observed that overall subjects were quite accurate when estimating brightness with the joystick, as revealed by the superposition of subjective
brightness estimations (green dots in the plots of Fig. 3) over the averaged joystick plots.

From the plots in Figure 3, it is clear that the time course of brightness perception is complex and that, except for one case, it differs substantially from the time course of stimulation. During these joystick experiments, we asked subjects to concentrate exclusively on brightness. However, this was a difficult task because they spontaneously and persistently reported that the size and color of percepts also changed during electrical stimulation. It thus appeared mandatory to complement brightness measurements with subjects' verbal reports describing the evolution of the quality (e.g., color and/or shape) of percepts. Table 2 summarizes subjects' descriptions. After analyzing all their comments, two general observations can be drawn. First, it is clear that only initial white/yellow percepts seem to be localized and bright enough to be used to construct a “useful” image. All subjects agreed on that statement. Second, past these initial instants, perception

<table>
<thead>
<tr>
<th>Subject</th>
<th>Joystick Plot</th>
<th>Verbal Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>S1</td>
<td><img src="image1" alt="Joystick Plot" /></td>
<td>Well-localized and bright percept of white color in the beginning followed by gradually decreasing brightness and becoming a very poorly defined blue “fat” line (“a light without shape”). Poorly localized and small reincrease in brightness at stimulus offset.</td>
</tr>
<tr>
<td>S2</td>
<td><img src="image2" alt="Joystick Plot" /></td>
<td>Brief (&lt;0.5 s), well-localized and bright percept of white/yellow color followed by an immediate decrease in brightness that changed rapidly to a “darker than background” percept. Poorly localized and medium reincrease in brightness at stimulus offset.</td>
</tr>
<tr>
<td>S3</td>
<td><img src="image3" alt="Joystick Plot" /></td>
<td>Well-localized and bright percept of yellow/orange color in the beginning, which after 2 s–3 s gradually decreases in brightness and “grows like an explosion” to fade into the “background.” Stimulus offset difficult to detect.</td>
</tr>
<tr>
<td>S4</td>
<td><img src="image4" alt="Joystick Plot" /></td>
<td>Well-localized and bright percept of white/yellow color remaining stable for about 5 s, which then disappears into the “background.” Well-localized and large reincrease in brightness at stimulus offset.</td>
</tr>
<tr>
<td>S5</td>
<td><img src="image5" alt="Joystick Plot" /></td>
<td>Well-localized and bright percept of yellow color in the beginning, fading into a “darker than background” percept at the end. “Background” at stimulus offset.</td>
</tr>
<tr>
<td>S6</td>
<td><img src="image6" alt="Joystick Plot" /></td>
<td>Well-localized and bright percept of white/yellow color that remains stable for the entire duration of the stimulus. At stimulus offset the percept changes to a blue light that fades into the “background.”</td>
</tr>
<tr>
<td>S7</td>
<td><img src="image7" alt="Joystick Plot" /></td>
<td>Brief (&lt;0.5 s), well-localized and bright percept of white color, immediately followed by a “dim reddish light” extending all over the visual field. Poorly localized and small reincrease in brightness at stimulus offset.</td>
</tr>
<tr>
<td>S8</td>
<td><img src="image8" alt="Joystick Plot" /></td>
<td>Well-localized and bright percept of white/silvery color in the beginning, followed by a dimmer orange light extending all over the visual field. Stimulus offset difficult to detect.</td>
</tr>
<tr>
<td>S9</td>
<td><img src="image9" alt="Joystick Plot" /></td>
<td>Brief (&lt;0.5 s), well-localized and bright percept of white/yellow color followed by a very dim “shimmering sensation” that disappears at stimulus offset.</td>
</tr>
</tbody>
</table>

The corresponding average joystick plots (see also Fig. 3) are included for comparison.
changed into what was most often described as dimmer and “shapeless” percepts covering large regions of the visual field and having different color. This second perceptual phase was qualified as much less useful (if useful at all) to reconstruct an image.

**Varying Stimulation Pulse Rate**

Figure 4 presents the averaged joystick plots (±SD) of each subject for a 10-second stimulus at 5 pps. Subjects S6, S8, and S9 reported similar joystick plots at this lower stimulation pulse rate than at 20 pps (compare with Fig. 3). For the remaining six subjects, lowering the stimulation pulse rate influenced the time course of brightness in different ways. For example, at 5 pps, S1 reported a substantially longer-duration percept (double the stimulus duration) than at 20 pps. In contrast, in the same stimulation condition, S3 reported a substantially shorter-duration percept than at 20 pps. Finally, at 5 pps, both “darker than background” percepts and reincreases in brightness observed at stimulus offset at 20 pps were practically suppressed.

Figure 5 presents the averaged joystick plots (±SD) of each subject for a 10-second stimulus at 60 pps. The joystick responses of subjects S6, S8, and S9 were similar to those obtained at the two lower stimulation pulse rates. For the remaining subjects, the effect of increasing the pulse rate was again variable. Subjects S3 and S4 reported substantially shorter-duration percepts at 60 pps than at 20 pps. Subjects S2 and S5 reported enhanced “darker than background” percepts. Finally, the 60-pps stimulation pulse rate tended to augment (or in some cases reveal) the brightness reincreases observed at stimulus offset. It is interesting to note that S4 reported that at 60 pps the brightness reincrease appearing at stimulus offset was considerably brighter than the initial flashlike percept appearing at stimulus onset.

**Varying Stimulus Duration**

Figure 6 presents the averaged joystick plots (±SD) of each subject for a 1-second stimulus at 20 pps. An interesting observation from this figure is that three of the nine tested subjects reported percepts that lasted longer than the stimulation. This was most striking for S1 and S8, where brighter than background percepts lasted as long as 10 seconds. At this shorter stimulus duration, S2 was the only subject to report a reincrease in brightness at stimulus offset.
Figure 7 presents the averaged joystick plots (±SD) of each subject for a 60-second stimulus at 20 pps. Five subjects (S1, S3, S7, S8, S9) reported percepts whose time course was similar to that observed at 10 seconds. For the remaining subjects, a few observations deserve to be highlighted. S2 described, after the initial flashlike and “darker than background” percepts, a brightness reincrease that disappeared beyond 30 seconds of stimulation. S5 described a “darker than background” percept after approximately 5 seconds, which remained fairly stable for the remainder of the stimulation. Subjects S4 and S6 reported that, after the initial stable percepts that lasted approximately 5 and 12 seconds, percepts disappeared completely for the remainder of the stimulation. It is interesting to note that S6, the only subject who reported the “ideal” time course of brightness for 10-second duration stimuli at 20 pps (i.e., a stable and bright percept lasting for the entire duration of stimulation), observed a fading percept beyond 12 seconds of ongoing electrical stimulation. In other words, for very long stimulation durations, this subject’s perception also had a dynamic and fading behavior, as observed for the other eight subjects. Finally, the brightness increases observed at stimulus offset were generally enhanced at this long stimulus duration.

Additional Experiments

Finally, in some subjects, we varied other parameters for control: stimulation amplitude (half and double the UCL), pulse width (3 ms per phase), testing the four single electrodes composing the tested QUAD separately, and testing an additional QUAD located as far as possible from the originally tested QUAD. When changing the stimulation amplitude to half or double the UCL, subjects described percepts as less/more bright in general but the time course of perceived brightness was similar (within experimental error). Percepts elicited by single electrodes were always reported as being smaller and less bright, but the time course of perceived brightness was essentially the same (within experimental error). As observed when varying stimulation pulse rate, we observed no general, systematic difference between the joystick plots obtained with a longer pulse width of 3 ms or when testing a different QUAD.

Discussion

Nine blind subjects using the Argus II Retinal Stimulation System participated in this study. They were asked to characterize their elementary visual perception on electrical stimulation of their retina. Of the nine tested subjects, only one
reported a well-localized, bright percept appearing at stimulus onset and lasting the entire duration of a 10-second stimulation trial. The others also reported well-localized and high brightness percepts at stimulus onset, but these percepts did not remain stable and well localized. Instead, they faded more or less rapidly, changing into different visual sensations that were described as being dimmer, poorly localized (covering large areas of the visual field), and having different color. Consequently, we can suppose that in everyday use of their retinal implant, these subjects are confronted with a difficult task: that of reconstructing images based on fading and changing percepts.

Intuitively, the amount of time during which precise visual information is available to subjects should have an impact on the visual performance that could be achieved with the device. In other words, not only should percepts be sharp and well localized, they should also last long enough for the brain to be able to reconstruct meaningful images. For example, it seems tremendously difficult to achieve accurate vision with flash-like percepts. Then, how much time should a well-localized and stable percept last for the brain to be capable of grasping the necessary information to reconstruct a patterned image? It is well known that in “normal” vision, visual information is exclusively gathered during fixations, except special situations (fixations are brief periods of time during which the eyes remain fairly stationary, between saccades). Normally sighted viewers have typical fixation durations of 200 to 250 ms during reading and of 260 to 330 ms during scene perception. The simple fact of restricting the number of characters visible at once (visual span) during normal reading significantly increases average fixation duration, and more than 400 to 500 ms are required for single character visual spans. Current electronic retinal prostheses provide very low resolution and a very limited “visual span.” Therefore, patients using these devices might require significantly longer “fixation” or “perceptual” times to grasp the necessary information. Indeed, we observed that in the visual tasks tested within the framework of the clinical trial, performance was generally poor for subjects where the duration of the initial, well-localized and high brightness percept was below 2 seconds. This was particularly true for tasks having the most stringent spatial vision requirements, such as character recognition (da Cruz L, et al. IOVS 2010; 51: ARVO E-Abstract 2023) and grating visual acuity. For example, the best score achieved to date in the grating visual acuity test (1.8 logMAR) was achieved by S6, the only subject for whom the initial well-localized percept lasted the entire duration of the 10-second stimulation trial. To our knowledge, none of the subjects...
participating in this study who experience flash-like percepts have been able to score reliably on this test (1.6–2.9 logMAR scale). We did not perform statistical analyses against performance data given the limited dataset available; however, this observation suggests a minimum percept duration to make practical use of the Argus II retinal implant.

One fundamental issue to be addressed is why electrical stimulation of the retina in human subjects elicits such variable and dynamic visual percepts. Although the contribution of adaptation mechanisms at structures high along the visual pathway cannot be excluded, there is some evidence suggesting it might be related to the complexity of retinal circuitry. Retinal prosthesis development was based on the fact that bipolar and ganglion cells are relatively spared in RP and ARMD, making them good targets for electrical stimulation. We do not know which retinal cells are being primarily activated by electrical stimulation of the retina in our subjects, but primarily activating one type of cell or another could have a significant effect on the type/quality of the elicited percepts. On one hand, animal studies suggest that the best strategy to achieve good temporal resolution would be to activate ganglion cells directly and avoid indirect activation through the retinal network. On the other hand, it has been

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**Figure 7.** Averaged joystick responses (red solid plots) ±SD (red dotted plots) versus time to 60-s duration stimuli presented at 20 pps for nine subjects. Note that the time scale used in the plots is different from in the previous figures. Each plot was calculated on the basis of the last five consecutive trials in this condition. The gray dotted plot represents stimulus duration. The green dots in the plots correspond to verbal brightness estimations made at critical time points. Each panel represents data from a single subject. Results were computed on the basis of five consecutive trials per subject and per condition. The black solid reference line shows the duration of the stimulus.

**Figure 8.** Mean duration (s ± SEM) of the FWLHB phase per subject for 10-s duration stimuli at 5 pps (black bars), 20 pps (light gray bars), and 60 pps (dark gray bars). This value was calculated as the duration of the first interval during which the joystick response remained ≥7. Results were computed on the basis of 5 consecutive trials per subject and per condition. The black solid reference line shows the duration of the stimulus.
postulated that the activation of the inner retinal network might result in better \textit{spatial resolution} than the direct stimulation of ganglion cells.\textsuperscript{47} Once the best neural targets in severely degenerated retinas have been identified, selective stimulation methods should allow for a better general outcome across patients.

Another interesting observation to be highlighted is the variability observed in the results, within and across subjects. In a given condition, the time course of brightness perception described by subjects was considerably different from one to the other. In addition, varying stimulation pulse rate had very different effects in each subject. This nonsystematic behavior is very difficult to interpret. Therefore, to further explore these variations, we tried to analyze separately the initial well-localized, high-brightness percept described by subjects as “useful” to construct an image. We calculated the duration of the initial “stable” percept, which we called the First Well Localized High Brightness (FWLHB) phase, at the three stimulation pulse rates tested. The duration of the FWLHB phase was computed as the amount of time that the joystick response remained above a brightness level of 7. This brightness criterion is somewhat arbitrary; but subjects were consistent in reporting that perception became shapeless at lower brightness levels. Figure 8 compares the duration of the FWLHB percept for all nine subjects, at the three pulse rates tested. The effect of stimulation pulse rate on the duration of the FWLHB percepts was also very variable. Subjects S2, S5, S7, and S9 showed virtually identical results in all stimulation conditions. For the others, changing the stimulation pulse rate influenced the duration of the FWLHB percept in different ways. For example, subjects S1 and S6 had the longest FWLHB percept durations at 5 pps (note that in the case of S6 this results in a percept lasting approximately 3 seconds longer than the stimulation). The longest FWLHB percept durations were obtained at 20 pps for subjects S3 and S4, and at 60 pps for subject S8. One-way repeated measures analysis of variance confirmed that, overall, the stimulation pulse rate did not significantly influence the duration of the FWLHB phase (F\textsubscript{2,16} = 0.318, \( P = 0.73 \)). Yet, an interesting outcome of this analysis is that, for some subjects, there is an “optimum” stimulation pulse rate for obtaining the best FWLHB percept duration results.

What are the reasons underlying this large variability? We checked for possible correlations between the duration of the FWLHB phase and relevant patients’ data, such as age at implant and time blind before implant. Because of the heterogeneous distribution of the different cell populations across the retina,\textsuperscript{48} we also investigated correlations between the duration of the FWLHB phase and the eccentricity of the tested QUAD. None of these variables correlated with the duration of the FWLHB phase (see Table 3). All the previous nonsystematic observations go in line with concerns raised by experts in the field of retinal remodeling. In retinal diseases like RP, retinal circuits are progressively remodeled through ongoing neural death, cell migration, and rewiring, resulting in anomalous synapses.\textsuperscript{49–52} Furthermore, there is considerable variation in the progression of the disease and the remodeling process, depending on the different RP variations. If the retinal circuitry is significantly remodeled and in different ways for each subject, it is reasonable to assume that the perceptual response to electrical stimulation would also differ considerably. Indeed, it has been proposed that patients with some residual cone function might be better candidates for retinal prostheses, as the integrity of the inner retinal layers could be better preserved.\textsuperscript{53} In future studies, the relationship between the implanted patients’ particular phenotype-genotype and the nature of their perceptual response to electrical stimulation of the retina should be thoroughly investigated. In addition, other retinal degenerations suitable for rehabilitation with a retinal prosthesis (e.g., ARMD) should also be considered.

\section*{Conclusion}

The perceptual response to electrical stimulation of the retina can be very different across subjects. Previous studies both in blind and normally sighted patients have already reported substantial differences in perception thresholds, shape/color of percepts, as well as performance.\textsuperscript{17–21,25,28,53,54} The present study demonstrates that the temporal properties of percepts evoked by electrical stimulation of the retina have a dynamic behavior that can vary substantially from subject to subject. Furthermore, only initial percepts at stimulation onset seemed to be useful to reconstruct a patterned image. Unfortunately, for several subjects, the duration of such initial percepts was very short.

Appropriate coding of a patterned image under such conditions appears challenging and will require careful selection of stimulation parameters. Significant research efforts are required to (1) understand how and why perceptual responses vary across patients, (2) determine the optimum stimulation strategies, and (3) if necessary, improve screening methods so that the candidates having the best rehabilitation prospects can be appropriately identified.

\section*{Acknowledgments}

The authors thank: Second Sight Medical Products, Inc. for facilitating this multicenter collaboration; the subjects for the time and energy they devoted into these experiments; Fatima Anafloos, Francesco Merlini, Brian Coley, Maura Arraraga-Chauvin, Celine Chaumette, Alexandre Leseigneur, Jeanne Haidar, Joe Zhong, and David Stewart for their collaboration; Greigore Cosendai for useful suggestions and discussions on the protocol and data analysis; and Daniel K. Freeman for helpful discussions on data from electrophysiology experiments.

\section*{References}


\begin{table}[h]
  \centering
  \caption{Simple (Pearson’s) Correlations of Relevant Patients’ and Performance Data versus the Duration of the FWLHB Phase for 10 s Duration Stimuli at 20 pps}
  \begin{tabular}{|l|c|c|}
    \hline
    \textbf{Age at implant} & \textit{R} = -0.34; \textit{P} = 0.36 & \\
    \textbf{Time blind before implant} & \textit{R} = 0.58; \textit{P} = 0.10 & \\
    \textbf{QUAD eccentricity} & \textit{R} = 0.09; \textit{P} = 0.80 & \\
    \hline
  \end{tabular}
\end{table}


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