Spatial extent of cochlear infrared neural stimulation determined by tone-on-light masking

Agnella Izzo Matic
Joseph T. Walsh, Jr.
Claus-Peter Richter
Spatial extent of cochlear infrared neural stimulation determined by tone-on-light masking

Agnella Izzo Matic, Joseph T. Walsh, Jr., and Claus-Peter Richter

Abstract. Artificial neural stimulation is widely used in clinic, rehabilitation, and research. One of the limitations of electrical stimulation is the current spread in tissue. Recently, pulsed mid-infrared laser stimulation of nerves has been investigated as an alternative stimulation method. The likely benefits of infrared neural stimulation (INS) include spatial selectivity of stimulation, noncontact mode of operation, and the lack of stimulation artifact in simultaneous electrical recordings. The hypothesis for this study is that INS of the cochlear spiral ganglion at low pulse energy is as spatially selective as low-level tonal stimulation of the cochlea. Spatial selectivity was measured using a masking method. An optical pulse with fixed optical parameters was delivered through a 200-μm diameter optical fiber. An acoustic tone, variable in frequency and level, was presented simultaneously with the optical pulse. Tone-on-light masking in gerbils revealed tuning curves with best frequencies between 5.3 and 11.4 kHz. The width of the tone-on-light tuning curves was similar to the width of tone-on-tone tuning curves. The results indicate that the spatial area of INS in the gerbil cochlea is similar to the cochlear area excited by a low level acoustic tone, showing promising results for future use of INS in implantable cochlear prostheses.

Keywords: pulsed diode laser; tuning curve; optical stimulation; cochlear implant; gerbil.

Introduction

A recent technological innovation provides focused neural stimulation using optical radiation. It has been demonstrated that auditory neurons can be stimulated with infrared radiation. This type of stimulation has been termed infrared neural stimulation (INS) and does not require the transgenic induction or exogenous application of chromophores. The use of lasers to evoke neural responses has several appealing features when compared to electrical stimulation: no direct contact is necessary between the stimulating source and the tissue (i.e., INS can be delivered through air), a finer spatial resolution of stimulation can be achieved, and no stimulation artifact is generated to deter simultaneous recordings of electrical responses from the neurons.

At many of the wavelengths used for optical stimulation, the radiation can be easily coupled to an optical fiber. Appropriate laser parameters will significantly affect the success of stimulation; e.g., wavelength dictates light scatter and absorption in the tissue. During INS, the stimulated area is confined to only the tissue along the optical path. Therefore, more spatially discrete populations of neurons should be stimulated as compared to electrical stimulation.

Previous work has demonstrated that optical stimulation is spatially selective in motor nerves and the cochlea. For the sciatic nerve, single fascicles could be activated during irradiation of the nerve trunk. The selectivity of optical stimulation in the cochlea was studied with an immunohistochemical method. Results showed more spatially confined areas of neural stimulation with INS as compared to electrical current. However, the method was biased because staining could only be achieved after exposure to high-level stimuli, tone pips, electrical current, or optical radiation. High-level stimuli likely activated larger populations of spiral ganglion cells, leading to a poorer spatial selectivity of stimulation. Therefore, to fully understand the benefits and limitations of optical stimulation, it would be important to determine the selectivity for low-level optical stimuli as well.

In this manuscript, spatial selectivity of cochlear INS has been measured using a real-time electrophysiological masking method. Masking refers to the ability of a masker stimulus (presented simultaneously or in short sequence with a probe stimulus) to reduce the response evoked by the probe stimulus alone. Dallos and Cheatham have, for example, shown that the cochlear compound action potential (CAP) amplitude evoked by a pure tone (probe) could be reduced by another pure tone (masker). At the level of the single neuron, simultaneous masking is manifested by a suppression of neural firing evoked by the probe alone. Resulting masker level-frequency curves (masker tuning curves or MTCs) showed a tuned response with a minimum at the frequency of the probe stimulus. Tone-on-tone MTCs reflect the activation of a local population of neurons along the cochlea, when using low-to-moderate probe tone levels. Thus,
the MTCs provide a measure of cochlear frequency selectivity at the site of stimulation. The recordings occur directly at the site of stimulation and do not reflect any upstream processing that may be reflected in neural responses recorded from the brain.

Here, “tone-on-light” MTCs were generated using a probe response evoked by INS and masked by a continuous acoustic masker. These experiments provide electrophysiological measurements that determine the cochlear region(s) stimulated by the laser. Since the cochlea has a tonotopic arrangement (high frequencies are encoded at the cochlear base, low frequencies at the cochlear apex), the physical location of the optical fiber along the cochlea can be translated into a corresponding frequency location of stimulation. Tone-on-light MTCs measure the spread of excitation during INS and can be compared with tone-on-tone MTCs.

2 Methods
All measurements were made in vivo using adult gerbils (*Meriones unguiculatus*). Animal care and use in this study was carried out in accordance with the NIH guidelines for the Care and Use of Laboratory Animals and was approved by the Northwestern University Animal Care and Use Committee. Eight animals were used for tone-on-tone masking experiments, and seven animals were used for tone-on-tone masking experiments. In some instances, multiple data sets, for both tone-on-light MTCs and tone-on-tone MTCs, were recorded in one animal.

2.1 Surgery
Animal surgery was performed as described previously. Briefly, gerbils were anesthetized by an initial intraperitoneal injection of sodium pentobarbital (80 mg/kg body weight). Maintenance doses were 17 mg/kg body weight and were given throughout an experiment whenever the animal showed signs of increasing arousal. Depth of anesthesia was assessed every 30 min by a paw withdrawal reflex. After the animal was fully anesthetized, breathing was facilitated by performing a tracheotomy and securing a length of PE90 tubing into the opening of the trachea. To maintain body temperature at 38°C, the animal was placed on a heating pad. Its head was stabilized in a heated head holder.

The cochlea was accessed using a ventral approach. The animal was positioned belly up and a dermal incision was made from the lower right jaw to the right shoulder in order to expose the right submandibular gland, which was subsequently ligated and removed. The muscles attached to the bulla and to the styloid bone were carefully dissected. Next, the bulla was opened, allowing access to the cochlea. A silver recording electrode was hooked onto the bony rim of the round window of the cochlea, and a ground electrode was placed under the skin at the left jaw. After cutting the cartilaginous outer ear canal, a speculum (probe) was fixed for each experiment. The acoustic masker was then plotted versus the masker frequency. A series of auditory nerve CAP threshold curves were obtained across frequencies in order to determine baseline cochlear function and verify a normal hearing animal. Acoustically evoked compound action potentials were measured using a modified tracking procedure. CAPs were measured between 2 and 50 kHz, with a resolution of 6 steps/octave. The acoustic stimuli were 12 ms tone bursts (including 1 ms rise/fall), which were presented in opposite phase. Thirty-two cochlear responses to the tone bursts were averaged for a single measurement. CAP threshold was defined as 30 ± 3 μV (N1/P1 amplitude). Acoustic CAP threshold curves that were acquired after the tone-on-light masking experiments showed no systematic shift in threshold, indicating that the cochlear preparation remained stable throughout the length of the experiment.

2.2 Monitoring Cochlear Function
Voltage commands for acoustic stimuli were generated using a computer I/O board (KPCI 3110, Keithley) inserted into a PC and were used to drive the speaker. The sound pressure out of the speaker was calibrated with a 1/8 in. Bruel and Kjaer microphone. All data are presented as calibrated sound pressure level in decibels (dB).

2.2.1 Compound action potentials
A series of auditory nerve CAP threshold curves were obtained across frequencies in order to determine baseline cochlear function and verify a normal hearing animal. Acoustically evoked compound action potentials were measured using a modified tracking procedure. CAPs were measured between 2 and 50 kHz, with a resolution of 6 steps/octave. The acoustic stimuli were 12 ms tone bursts (including 1 ms rise/fall), which were presented in opposite phase. Thirty-two cochlear responses to the tone bursts were averaged for a single measurement. CAP threshold was defined as 30 ± 3 μV (N1/P1 amplitude). Acoustic CAP threshold curves that were acquired after the tone-on-light masking experiments showed no systematic shift in threshold, indicating that the cochlear preparation remained stable throughout the length of the experiment.

2.3 Masking Procedures

2.3.1 Tone-on-tone masking
Stimuli consisted of a probe tone and a masker tone. The probe tone (12 ms duration, including a 1 ms rise/fall) was simultaneously overlapped by a masker tone (34 ms duration including a 5 ms rise/fall). Stimuli were presented at a rate of 4 Hz. Tone-on-tone masking data were acquired using CAPs.

To measure tone-on-tone MTCs, the frequency of the probe tone was fixed and was presented at a given sound level such that the probe-evoked CAP was approximately 40 to 50 μV. The data were averaged for 32 presentations of the tone.

2.3.2 Tone-on-light masking
Masking experiments were conducted with laser pulses as the probe stimulus, and continuous acoustic tones as the masker. The placement of the optical fiber in the cochlea determined the “frequency” of stimulation. The energy level of the laser (probe) was fixed for each experiment. The acoustic masker was varied in frequency and in level, as described previously for tone-on-tone masking experiments.

A pulsed diode laser (wavelength = 1.86 μm, pulse duration = 20 μs, repetition rate = 2 Hz; Research Infrared Nerve Stimulator, Lockheed-Martin Acuity Corp.) was used to generate the probe stimulus. Previous work has examined the wavelength variability of INS in the sciatic nerve and in the cochlea.

The current wavelength was selected based on data that showed an optimal margin of safety between stimulation and ablation.
in the sciatic nerve at wavelengths with a similar penetration depth. In contrast to earlier studies that used a shorter penetrating wavelength that required the optical fiber to be placed at the modiolus to evoke cochlear responses, the wavelength used in this study had a longer penetration depth and, therefore, the optical fiber could be placed such that the tip was approximately 0.5 mm from the spiral ganglion cells and was in contact with the fluids on the round window membrane.

The laser was coupled to a 200-μm core diameter optical fiber (FIP series, Polymicro), which was inserted through the opening in the bulla and abutted to the round window membrane (Fig. 1). The optical fiber was mounted on an x-y-z translator (MMW-203, Narishge) attached to a micromanipulator to control the fine position. The optical fiber was in contact with the round window membrane, but did not penetrate it. The fiber was visually oriented toward the spiral ganglion cells in Rosenthal’s canal in the upper basal turn, which is approximately 2.5 mm along the length of the basilar membrane from the most basal end of the cochlea. At the wavelength used here, the ~10 μm thick round window membrane likely has a negligible effect on the optical beam and can be modeled by water absorption, which the optical beam and can be modeled by water absorption, forward scattering, and multiple reflections. When irradiating the spiral ganglion through the round window membrane, but did not puncture it. The orientation was toward the lateral position of the round window (left in this picture). Various tuning curves were acquired with the optical fiber in different positions in the round window along the trajectory from lateral to medial (left to right). The fiber was mounted on an x-y-z translator coupled with a micromanipulator. (See also Ref. 18 for optical fiber placement.)

2.4 Data Analysis

Resultant tone-on-tone and tone-on-light MTCs were analyzed with Igor Pro software (WaveMetrics). Each data set was interpolated to 100 points using a smoothing cubic spline function based on the nearest 3 points. All measurements were made on the interpolated data.

For tone-on-tone masking paradigms, the frequency at which the MTC shows the lowest masker sound level is termed the best frequency (BF). The BF of the tuning curve corresponds to the frequency of the probe tone (Fig. 2). The sharpness of tuning describes the selectivity of the neural response. One method of characterizing the sharpness of tuning at the BF is by the bandwidth (BW) of the tuning curve at 10 dB above the MTC tip level. The sharpness of tuning (Q10dB) was calculated as follows:

\[
Q_{10\text{dB}} = \frac{\text{BW}}{\text{BF}}
\]

(see Fig. 3 for a graphical representation of tuning curve calculations). In addition to sharpness of tuning, the slope of the high-frequency side of the MTCs was calculated. The slope (in dB/octave) was measured between 20 and 50 dB above the BF masker level on the high frequency side of the MTC. The tip-to-tail ratio was measured as the ratio between the masker level at the BF to the masker level at one octave below the BF. Statistical significance was calculated with a two-tailed Student’s t-test at \( \alpha = 0.5 \).

3 Results

A masking method was used to determine the spatial selectivity of acoustic and infrared neural stimulation in the gerbil cochlea. MTCs were measured and compared for an acoustic probe and for a laser probe. The resulting tone-on-tone MTCs exhibit a tuned response, with a characteristic tip at the best frequency and sloped sides (Fig. 4). The tone-on-light MTCs exhibit more variability in the shape of the MTC, as compared to the tone-on-tone MTCs. When irradiating the spiral ganglion through the round window, the optical beam diverges approximately 3 deg from collimated (unpublished). As previously shown, the exiting beam had a radial Gaussian profile. Radiant exposures at the tip of the fiber were between 10 and 20 mJ/cm² as measured by an energy sensor (J50LP-1A, Coherent-Molecron). The radiant energies evoked a CAP of 40 to 50 μV, moderately above threshold. The noise level of the recording system was ~6 μV and the INS-evoked CAP amplitude was typically 250 μV at saturation

An unmasked optically-evoked CAP was recorded to determine the baseline response. The responses to 10 stimulus presentations were averaged. Next, the continuous acoustic masker tone was presented in addition to the laser pulse. The acoustic masker was on for 2 to 3 s prior to the 10 repetitions of the laser pulse being presented. For each masker frequency and level, the INS-evoked CAP was measured. Similar to the tone-on-tone experiments, the masker level that was needed to decrease the INS-evoked CAP by 6 dB was recorded. Experiments began with the masker at 32 kHz, decreasing by 6 steps/octave, to 1 kHz. The sound level at each acoustic frequency began with the maximum output from the speaker and was decreased in steps of 5 dB until the criterion was reached, a decrease of the probe’s response by 6 dB. The resulting level of the masker was then plotted versus the corresponding frequency. A tuned frequency-level curve resulted. To document any changes in cochlear function due to laser irradiation, acoustic CAP threshold measurements were conducted at the conclusion of the experiments.
Fig. 2 Representative tone-on-tone MTC with calculations for data analysis. This tone-on-tone MTC shows the best frequency (tip of the tuning curve) and the sharpness of tuning, $Q_{10\text{dB}}$, which was determined by the ratio of BF to the MTC width at 10 dB sound level above BF level. For the tuning curve shown, BF = 10 kHz and the level at BF is 50 dB. At a sound level of 60 dB, or 10 dB above the BF level, the width of the curve extends from 9.4 kHz to 10.4 kHz, giving $Q_{10\text{dB}} = 10/(10.4 - 9.4) = 10$. The slope was measured as the rise/run between 20 and 50 dB above the BF level. When the high frequency side of a curve did not extend 50 dB above the BF level, we calculated the slope to the highest masker level. At 20 dB above the BF level, or 70 dB, the frequency is 11.2 kHz, and at 50 dB above the BF level, the frequency is 13.4 kHz. Therefore, the slope = $(100 - 70)/(13.4 kHz - 11.2 kHz) = 115.9 \text{dB/octave}$. The tip-to-tail ratio was measured as the ratio between the masker level at the BF to the masker level one octave below the BF. For this tuning curve, one octave below the BF occurs at 5 kHz, however, the data do not extend to quite this low of a frequency. When the low frequency tail did not extend one full octave below the best frequency, as is the case with this curve, the masker level at the lowest frequency of the tail was used. At this frequency, we measured a sound level of 97 dB, giving a tip-to-tail ratio $= 50/97 = 0.51$.

Fig. 3 Tone-on-light MTCs demonstrate spatially selective infrared neural stimulation. Several examples of tone-on-light MTCs, shown in solid lines, are compared with the tone-on-tone MTCs, shown in dotted lines. The data are presented on three separate graphs, grouped according to BF, for clarity of presentation. Each tone-on-tone MTC shows the raw data measured from one trial in one animal with a fixed optical fiber position and fixed laser energy/pulse. Each tone-on-tone MTC shows the raw data measured from one masking trial in one animal at one fixed probe frequency and level. Optical fiber locations were varied between trials, therefore leading to MTCs with different BFs. Both optical and acoustic probe levels were selected that a 40 to 50 $\mu$V CAP resulted when the probe was presented alone.

Matic, Walsh, and Richter: Spatial extent of cochlear infrared neural stimulation...
light MTCs were not statistically significantly different from ratios from tone-on-tone MTCs ($p=0.07$).

When the optical fiber was moved from a more apical irradiation orientation to a more basal irradiation one, the resulting tone-on-light MTCs had different BFs. Figure 7 shows one example of that behavior. The BFs of these curves were 5.3 and 10.8 kHz, which is in close agreement with the amount of frequency change expected for a $\sim 1$ mm translation, based on a 1.4 mm/octave frequency-place slope of the gerbil. In other animals, when the optical fiber was translated 200 to 500 $\mu$m within the round window, MTCs shifted BFs from 9.4 to 8.3 kHz and from 8.6 to 6.1 kHz.

### 4 Discussion

To assess the spatial spread of optical stimulation in the gerbil cochlea, tone-on-light masking experiments were conducted. Here, the response to a laser pulse was masked with continuous acoustic tones. The resulting tone-on-light MTCs were qualitatively and quantitatively similar to the tone-on-tone MTCs. Tone-on-light MTCs had BFs between 5.3 and 11.4 kHz. The BFs correspond to the irradiation location of optical fiber along the spiral ganglion. Previous experiments using the same optical fiber orientation delivering high-level optical stimuli have shown neural activation in the upper basal portion of the cochlea, about one turn above the tip of the optical fiber. In addition, the beam path of INS was measured in the guinea pig cochlea. These
data demonstrated that, due to the surgical access obtained and the orientation of the optical fiber, it was extremely difficult to target neurons in the very base of the cochlea. Most often, neurons 1/4 to 1 full turn more apical from the surgical access were targeted.

Given that the length of the gerbil basilar membrane is ~11.4 mm in total and the hearing range of the gerbil is between 0.14 and 63 kHz, the gerbil frequency-place map has a slope of ~1.4 mm/octave. Hence, the frequency of the neurons opposite to the optical fiber in the upper basal turn of the gerbil cochlea should be centered around the 7 kHz location. Indeed, the range of BFs measured for the tone-on-light MTCs, from 5.3 to 11.4 kHz, include this calculated location. However, keep in mind that 7 kHz is an approximate location because the above data are calculated based on the frequency-place map of the basilar membrane, and not for the spiral ganglion, the likely location of INS depolarization. It is known that the spiral ganglion can have a compressed and/or shifted frequency-place map relative to that of the basilar membrane. The corresponding change in best frequency, as shown in Fig. 1, demonstrates that the placement of the optical fiber determines the location of stimulation in the cochlea and thus the frequency.

The most important information that can be gained from the tone-on-light MTCs is the spatial confinement of INS. Here, the measure of the spatial spread of stimulation is expressed by the $Q_{10dB}$ ratio. With increasing $Q_{10dB}$ values, the stimulated area decreases. The tone-on-light $Q_{10dB}$ ratios were statistically the same as the tone-on-tone $Q_{10dB}$ ratios. Therefore, with a 200-μm diameter optical fiber, the results suggest that the areas of optical and acoustic neural stimulation are approximately the same at low stimulus levels. It is likely that a decrease in the optical fiber diameter would decrease the area of neural stimulation, which would be reflected as a narrower tuning curve (higher $Q_{10dB}$). Data measuring the spatial extent of cochlear INS using inferior colliculus recordings also indicated spatially restricted areas of stimulation with similar BFs. The advantage of the inferior colliculus (IC) recordings is that they can be achieved in an acutely deafened animal model and multiple locations in the IC can be probed simultaneously. These data, however, were recorded from the auditory midbrain and potentially reflect processing of the neural signals upstream from the cochlea. The experiments presented here are a direct measurement of the evoked cochlear responses at the site of stimulation.

Tone-on-tone MTCs measured in this study are comparable to published data. $Q_{10dB}$ values for acoustic MTCs previously recorded from the gerbil CAP ranged from ~2 to 7, which overlaps with the range of values reported here. It has been shown that continuous acoustic MTCs and simultaneous pulsed acoustic MTCs show similar sharpness of tuning. Therefore, the data collected with continuous acoustic masking of pulsed optical radiation can be compared to simultaneous pulsed tone-on-tone MTCs.

INS is likely mediated by a photothermal mechanism. The radiation is primarily absorbed by water in the tissue and this energy is dissipated as heat. Recent evidence indicates that secondary to the transient temperature rise, a change in the capacitance of the lipid occurs. However, the presence of a temperature-dependent capacitance change does not preclude the existence of additional mechanisms mediating INS. It is unlikely that a primary photomechanical mechanism causes the neural depolarization given that the optical parameters here operate outside of stress confinement. Nevertheless, we have examined INS in acute and chronic deafened animals and shown that they do not have significantly elevated optical stimulation thresholds at short pulse durations when compared to normal hearing animals. In the chronic deafened animals, there were no remaining hair cells, negating the possibility that INS in the cochlea is mediated by a laser-induced pressure wave in the perilymph. However, there were remaining spiral ganglion neurons, the likely location for INS action. For the experiments presented here, deafening the animals would have abolished the ability to achieve acoustic masking and rendered the experiments unfeasible.

One might speculate on the benefits of this method of neural stimulation for neural interfaces, in particular cochlear implants. Cochlear implants are one of the most successful clinical neuroprostheses, restoring sound perception to severely deaf individuals by electrically stimulating auditory neurons in the inner ear. However, electrical stimulation can suffer from electric current spread in tissue, preventing spatially localized stimulation of the neural tissue. It has been argued that the performance of neural interfaces, including cochlear implants, could be improved significantly if more discrete locations of neurons could be stimulated simultaneously. Various modifications to traditional electrical stimulation paradigms are under investigation to increase the spatial selectivity of stimulation and the performance of cochlear implant users. A more spatially selective stimulation paradigm using INS may also allow for more
independent stimulation channels to be presented simultaneously. Here, we demonstrated that cochlear INS activates a select-ive portion of the cochlea, on the order of a low level tone.

Before designing an optical cochlear implant, though, studies on chronic stimulation and safety of INS will be conducted, particularly at high rates of stimulation that are relevant to the cochlea and cochlear implants. Acute studies of high repetition rate stimulation demonstrate no change in CAP amplitude while stimulating up to 200 Hz for up to 8 h. Other parameters, such as placement of optical sources, will be examined and optimized for an optical cochlear implant. In summary, the results provided in this paper support the view that optical radiation can provide spatial selectivity of stimulation similar to acoustic stimulation of the cochlea. Optical radiation can be directed and confined to a small volume of tissue, can stimulate small populations of neurons, and can be deployed in a noncontact fashion (through air) or in a fluid interface, which leads to little divergence of the optical radiation beam.

Acknowledgments

This project has been funded in whole or in part with federal funds from the National Institute on Deafness and Other Communication Disorders, National Institutes of Health, Department of Health and Human Services, under Contract No. HHSN260-2006-00006-C/NHI No. N01-DC-6-0006, by NIH Grant Nos. R41 DC008515-01 and F31 DC008246-01, and the E. R. Capita Foundation. The authors would like to thank Acclivity Corporation for lending the laser that was used for these experiments.

References

