

EVALUATION OF IMPLANTED EPIRETINAL MICROCONTACTS IN THE MAMMALIAN RETINA* ((N.Peixoto¹, S.Sträßburger¹, R.Hornig¹, P.Walter², P.Szurman², R.Eckmiller¹)) 1) Dept. Computer Science VI, University Bonn; 2) Dept. Ophthalmology, Univ. Cologne, F.R. Germany.

Purpose. To explore the optimal electrical stimulation parameters for physiologically acceptable elicitation of neural responses from retinal ganglion cells. **Methods.** We developed a programmable stimulus pattern generator (PSPG) for generation of single- or multi-channel constant current pulses with variable biphasic shapes, pulse train duration, repetition rate, and varying degrees of synchronicity in case of multi-channel stimulation. Electrically evoked potentials (EEPs) from the cortex of anesthetized rabbits were recorded (see: Walter et al., ARVO, 1998). **Results.** Thresholds for EEPs were lower for trains of 5 to 10 biphasic pulses applied to several adjacent microcontacts simultaneously. Whereas single electrode stimulation required 40 to 50 μ A per phase at threshold, a measurably smaller amplitude was sufficient for simultaneous stimulation of 2 or 3 adjacent electrodes. A comparative analysis of EEP amplitudes revealed that biphasic stimulation with pulse trains even at the unphysiologically high level of 200 μ A did not damage the retinal tissue, whereas this is clearly the case for monophasic stimulation. **Conclusions.** Our constant current PSPG is a powerful and adequate device for stimulus parameter optimization in conjunction with implanted epiretinal microcontacts. EEP thresholds in the physiologically acceptable range far below 100 μ A can be achieved even with the non-penetrating microcontacts, which were developed by our partners (FhG-IBMT, FhG-IMS). Threshold reduction by means of simultaneous stimulation of several electrodes supports the hypothesis that supra-threshold EEPs represent inputs from a larger number of retinal ganglion cells and that accordingly the threshold for single ganglion cell stimulation is even lower.

For details of the retina implant project see: <http://www.nero.uni-bonn.de/ri/retina-en.html>

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