

Supplementary Information

Model-based prediction of optogenetic sound encoding in the human cochlea by future optical cochlear implants

Lakshay Khurana^{1,2,3,4,5}, Daniel Keppeler^{1,3,5}, Lukasz Jablonski^{1,2,5,#}, and Tobias Moser^{1,2,3,5,6,#}

¹Institute for Auditory Neuroscience, University Medical Center Göttingen, Göttingen, Germany

²Auditory Neuroscience and Optogenetics Laboratory, German Primate Center, Göttingen, Germany

³Auditory Neuroscience & Synaptic Nanophysiology Group, Max Planck Institute for Multidisciplinary Sciences, Göttingen, Germany

⁴Göttingen Graduate Center for Neurosciences, Biophysics, and Molecular Biosciences (GGNB), University of Göttingen, Göttingen, Germany

⁵InnerEarLab, University Medical Center Göttingen, Göttingen, Germany

⁶Cluster of Excellence "Multiscale Bioimaging: from Molecular Machines to Networks of Excitable Cells" (MBExC), University of Göttingen, Göttingen, Germany

#correspondence: lukasz.jablonski@wp.eu, tmoser@gwdg.de

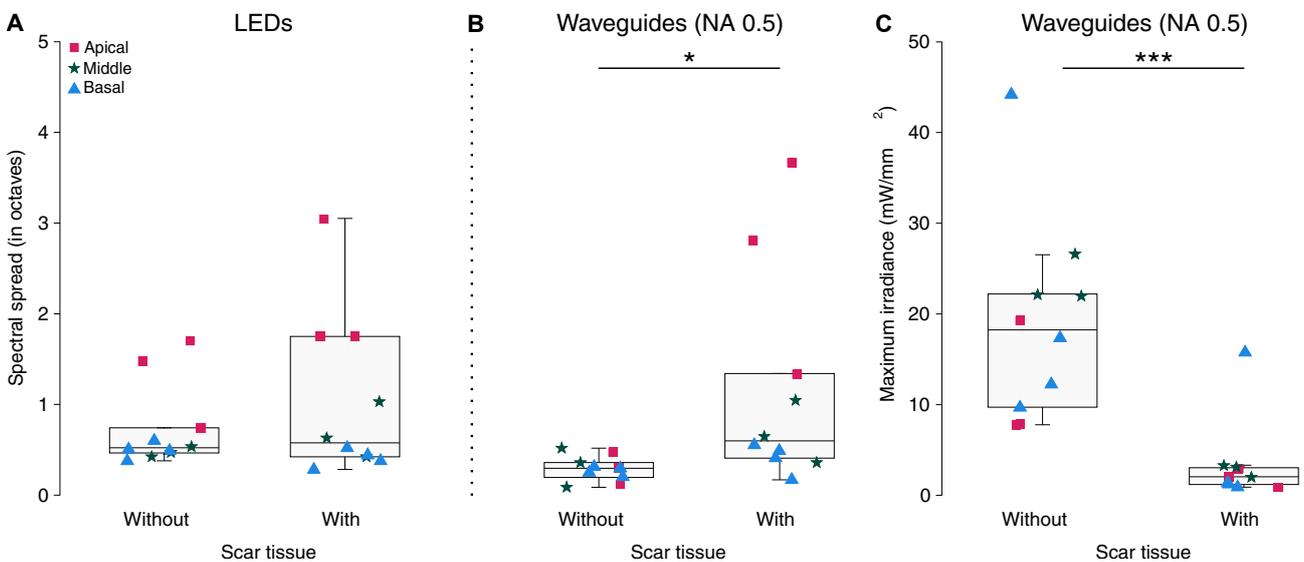


Figure S1: Effect of fibrosis on irradiance and spectral spread. (A) Impact of scar tissue on spectral spread for LED emitters when the model was simulated with scala filled with scar tissue. The whiskers extend to the maximum and minimum data points not considered outliers. **(B)** Impact of scar tissue on spectral spread for the waveguide emitters. **(C)** Decrease of maximum irradiance values for waveguide emitters (NA 0.5) when the model was simulated with scala filled with scar tissue. Box plots indicate median (center line), as well as 25th percentile and 75th percentile as the bottom and top edges. The asterisks represent statistical significance (* indicates $p < 0.05$, ** indicates $p < 0.01$, and *** indicates $p < 0.0001$).