

Table S1. Nucleotide sequences of primers used for PCR amplification

Gene	Forward Primer (5' to 3')	Reverse Primer (5' to 3')
<i>Bmp2</i>	ATCCTGAGCGAATTCGAGTTG	GGGTGGGTCTCTGCTTCAAG
<i>Id3</i>	AACAGAGCCTTTCTCCAAGGAA	TGTCCGAACTCTGCCAAGGT
<i>Nog</i>	GGTCGAAGATAGGGTCCAAGTG	TCCAATTCCCAGCGACAAC
<i>Gapdh</i>	GCAACCCGAGACAAGATGGT	GCGTCCAATACGG CCAAAT

Bmp2, bone morphogenetic protein 2; *Id3*, inhibitor of DNA binding 3; *Nog*, noggin; *Gapdh*, glyceraldehyde-3-phosphate dehydrogenase.

Supplementary Materials Figures

Figure S1

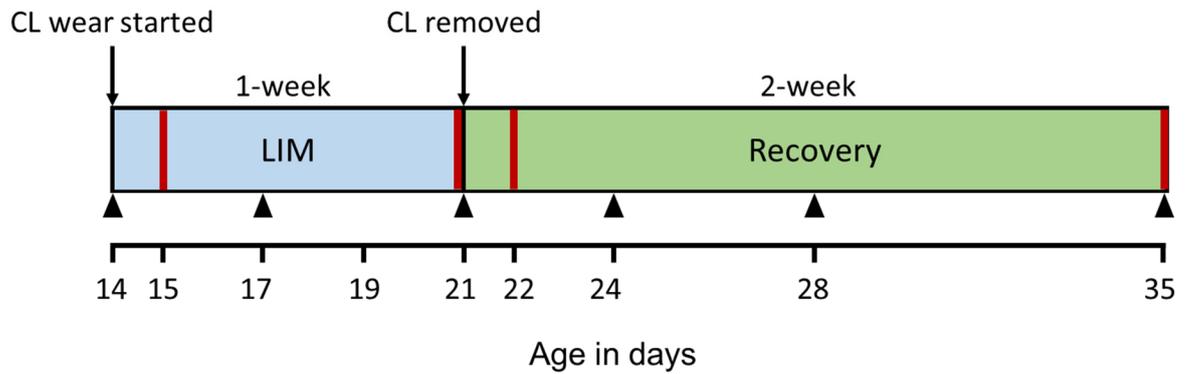


Figure S1. Study design, showing the contact lens (-10 D) wearing schedule, along with the time lines for *in vivo* measurements and collection of retinal pigment epithelium (RPE) samples for mRNA isolation. The black arrowheads indicate the time points at which refractive error, axial length, as well as choroidal, and scleral thickness data were collected; the red vertical bars indicate the time points at which RPE samples were collected. LIM: lens-induced myopia.

Supplementary Materials Figures

Figure S2

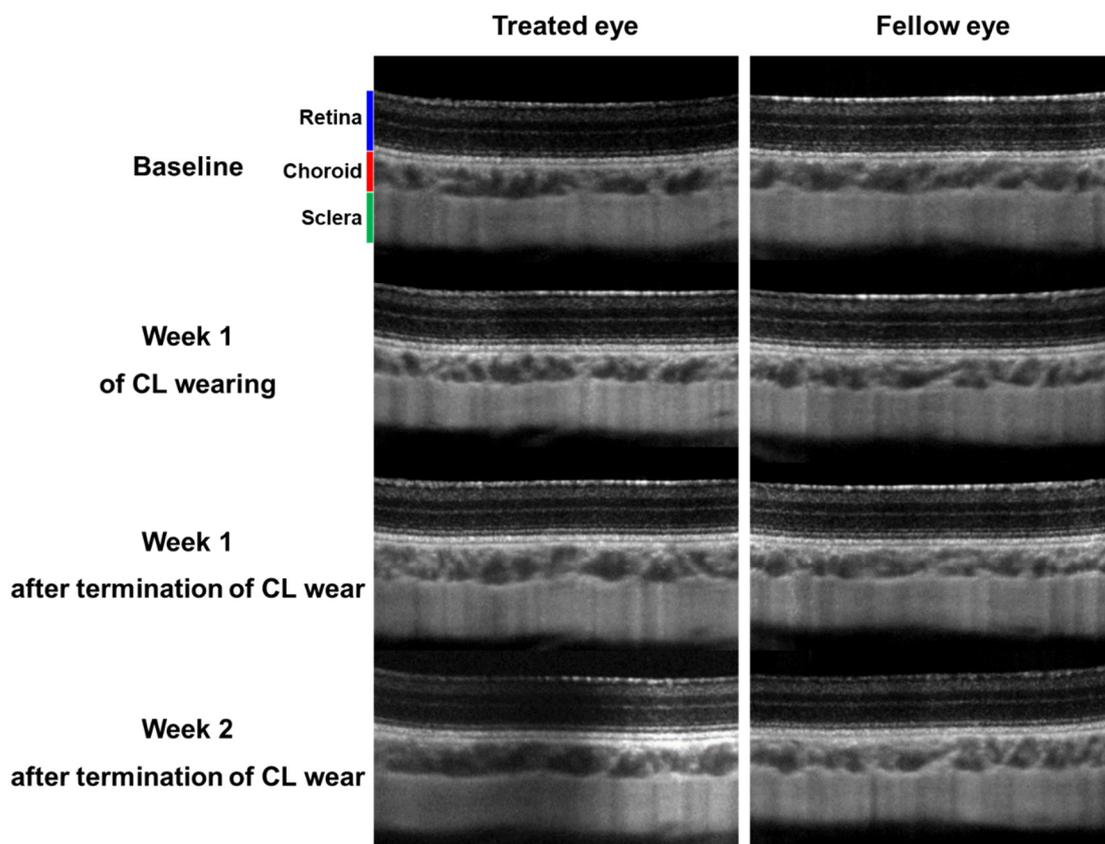


Figure S2: Representative SD-OCT fundus images from treated and fellow eyes of the same guinea pig captured 1 mm above the optic nerve head, on day 0 (pretreatment baseline), after 1 week of CL wear (-10 diopter), and week 1 and week 2 after termination of CL wear. Retinal, choroidal, and scleral boundaries are clearly visible in all images.