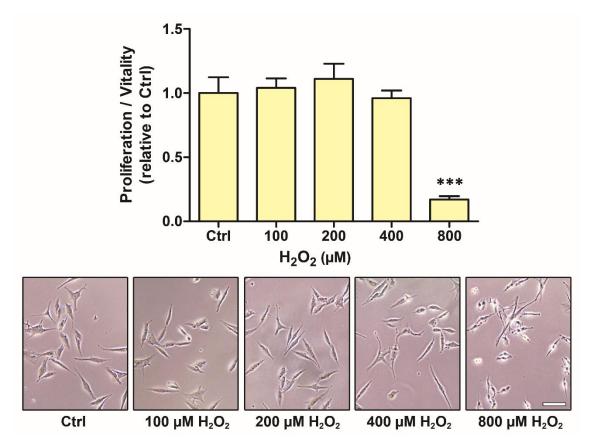
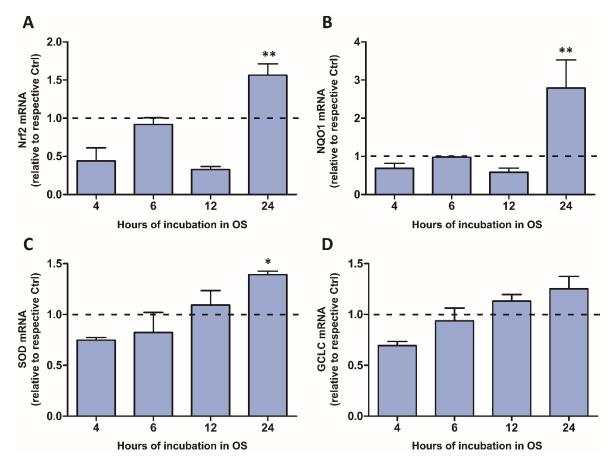


**Figure S1.** Morphology of living MIO-M1 cells (**A**) and of MIO-M1 cells after fixation for 20 min in 4% paraformaldehyde in PBS and permeabilization with 0.3% Triton X-100 in PBS for 5 min (**B**). DIC microscopy. Scale bar, 100 μm.



**Figure S2.** Proliferation/vitality of MIO-M1 cells treated with different doses of H<sub>2</sub>O<sub>2</sub> to determine the H<sub>2</sub>O<sub>2</sub> concentration to be used in the experiments with MIO-M1 cells. \*\*\* p < 0.001 relative to controls (Ctrl). n = 4. The photomicrographs were taken with DIC microscopy. Scale bar, 100 µm.



**Figure S3.** Time course of *Nrf2* and of oxidative stress marker expressions in response to OS to determine the shortest incubation time for detecting an effect of OS in retinal explants. \* p < 0.05, and \*\* p < 0.01 relative to the respective control, represented by the dashed line. Two-way ANOVA followed by Bonferroni post-hoc test. n = 3.