Supplementary Materials: NLRP3 Upregulation in Retinal Pigment Epithelium in Age-Related Macular Degeneration

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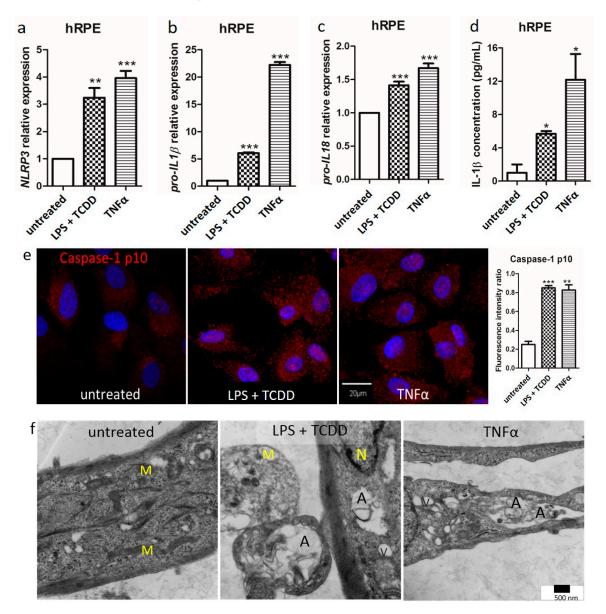


Figure S1. Activation of NLRP3 inflammasome in primary adult human RPE (hRPE) cells under inflammation and oxidative stress. (**a**–**c**) qRT-PCR analysis of *NLRP3*, pro-*IL1* β and pro-*IL18* in hRPE cells stimulated with LPS + TCDD and TNF α (n = 4); (**d**) ELISA analysis of IL-1 β in the supernatants of hRPE cells treated with LPS + TCDD and TNF α (n = 4); (**e**) Confocal microscopy of hRPE stimulated for 24 h with LPS + TCDD and TNF α (n = 4). Caspase-1 p10 subunit was labeled in red. The nuclei were stained with 4,6-diamidino-2-phenylindole dihydrochloride (DAPI) (blue). Image-J software is used to measure the fluorescence intensity in pixels per area in each image and expressed as fluorescence intensity ratio. Scale bar = 20 μm; (**f**) Ultrastructural evaluation of hRPE cells stimulated with LPS + TCDD and TNF α . Scale bar = 500 nm. N, nuclear; M, mitochondrion; A, autophagosome; V, vesicle. Data are presented as mean ± SEM. * p < 0.05, ** p < 0.01. *** p < 0.001.

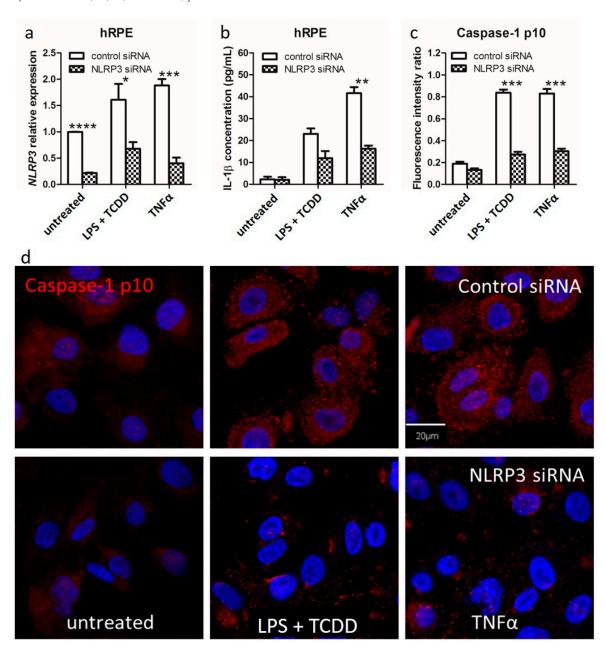


Figure S2. NLRP3 knockdown inhibits inflammasome activation in hRPE cells under inflammation and oxidative stress. (**a**) qRT-PCR analysis of *NLRP3* in hRPE cells stimulated with LPS + TCDD and TNF α for 24 h after siRNA transfection (n = 4); (**b**) ELISA analysis of IL-1 β in the supernatants of hRPE cells treated with LPS + TCDD and TNF α after siRNA transfection (n = 4); (**c**) The Caspase-1 p10 protein ratio of immunohistochemistry is calculated. Image-J software is used to measure the band intensity in pixels (n = 4); (**d**) Confocal microscopy of hRPE cells treated with LPS + TCDD and TNF α for 24 h after siRNA transfection (n = 4). Caspase-1 p10 subunit was labeled in red. The nuclei were stained with DAPI (blue). Scale bar = 20 μm. Data are presented as mean ± SEM. * p < 0.05, *** p < 0.01, **** p < 0.001.