

Figure S2. Quantification of AFGs in pRPE. pRPE were pulsed once with POS or UV-irradiated POS and chased for the indicated time points before fixing and imaging AF in the 488 emission channel with a high-content imaging platform. (A) Representative maximum intensity projections of confocal slices are shown. Blue: DAPI; Green: AF. Scale bar: 15 μm (B) The difference in the number of granules per cell between cells with no POS and POS, at the different time points, and the number of granules per cell overtime in (C) cells incubated with POS or (D) UV-irradiated POS (UVPOS) were quantitated from the z-stacks. Values are represented as mean ± SEM of the number of AF punctae/number of DAPI punctae per field of view. A single biological replicate with a minimum of 1500 cells from at least 4 wells was analysed per condition. Statistical differences, relative to cells with no POS, were performed using the one one-way ANOVA followed by Kruskal Wallis test and differences relative to POS 0h were performed using Mann-Whitney t test. (\*\*\*\*\*P<0.001).