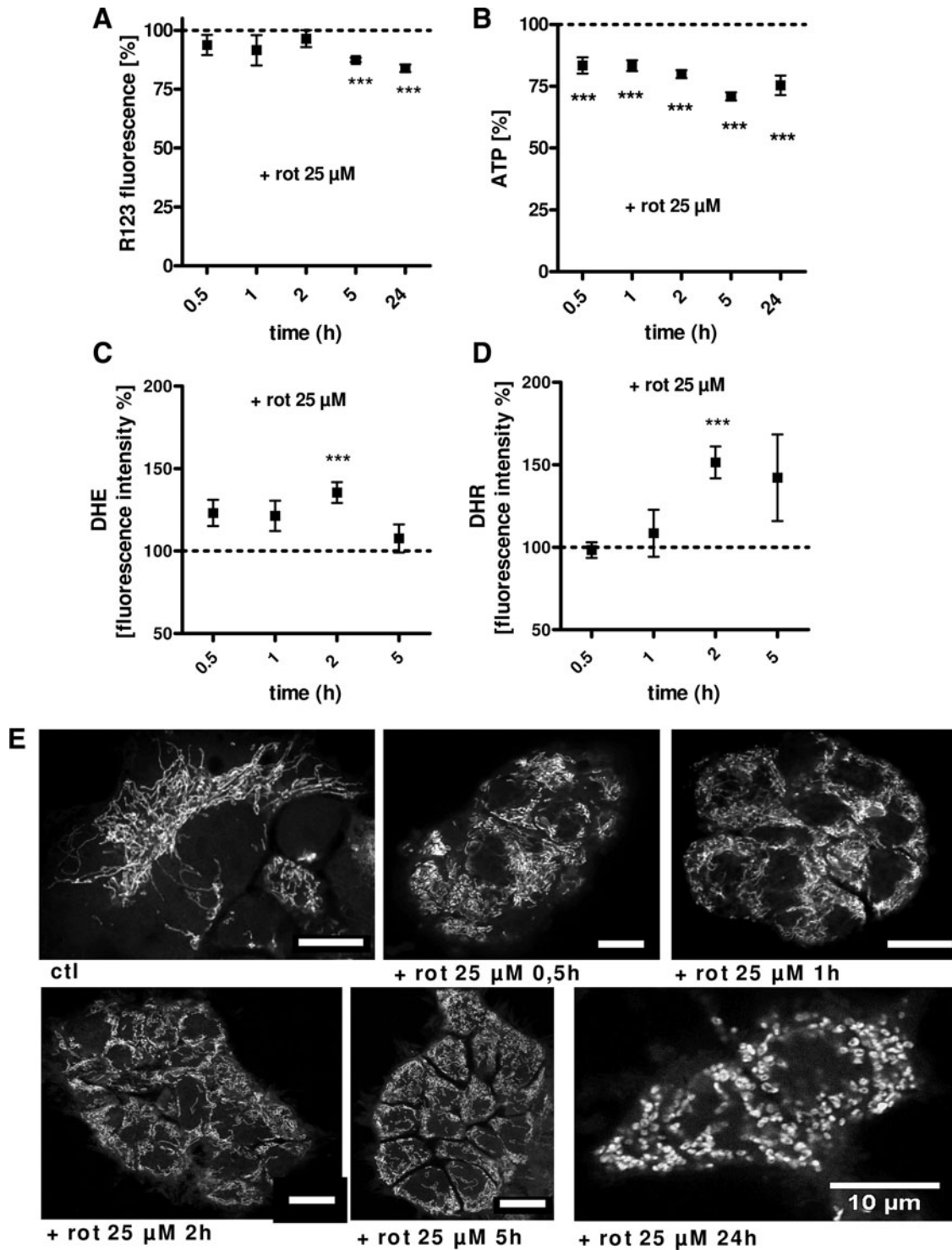
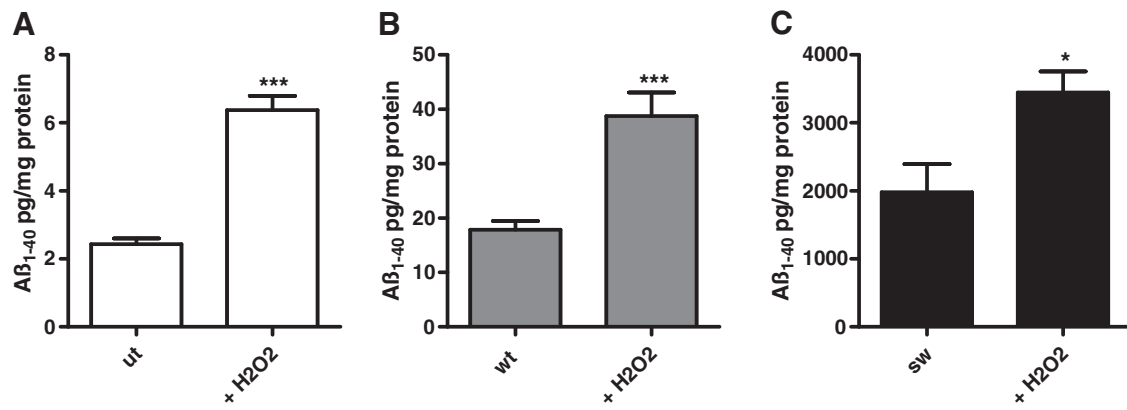


Supplementary Data



SUPPLEMENTARY FIG. S1. Time-dependent effect of rotenone (rot) 25 μ M on mitochondrial function in untransfected human embryonic kidney (HEKut) cells. (A) Time-dependent effect of rotenone (25 μ M) on mitochondrial membrane potential (R123 fluorescence intensity), ATP levels (B), superoxide anion radicals (dihydroethidium [DHE] fluorescence, C), and cytosolic reactive oxygen species (ROS; dihydrorhodamin [DHR] fluorescence, D) in HEKut cells. (E) Representative confocal images of the time-dependent effects of rotenone (25 μ M) on the mitochondrial morphology in HEKut cells (bars represent 10 μ m) compared to untreated controls after mitochondrial staining with MitoTracker CMXRos. (A–D) $n=6 \pm$ standard error of the mean (SEM); unpaired t -test; *** $p < 0.001$.



SUPPLEMENTARY FIG. S2. Oxidative stress induces A β generation. HEK ctrl (A), wild-type amyloid precursor protein (APPwt) (B), and APP containing the Swedish mutation (APPsw) (C) cells were treated for 24 h with H₂O₂ and soluble A β ₁₋₄₀ levels were investigated. (A–C) $n=4-6 \pm$ SEM; unpaired t -test; * $p < 0.05$, *** $p < 0.001$.