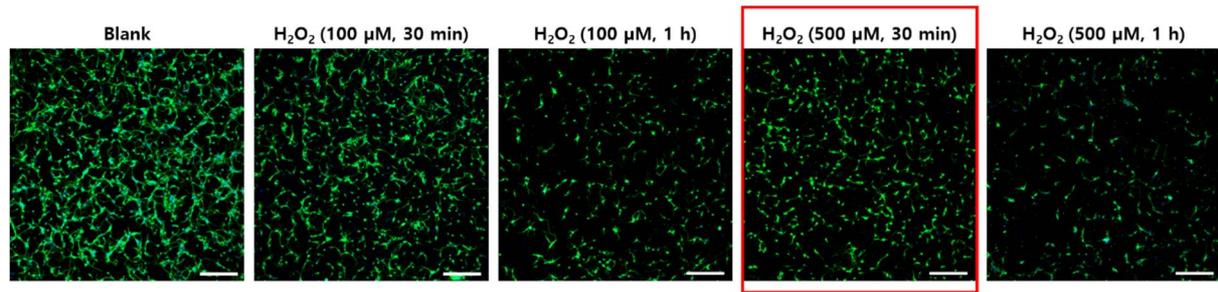
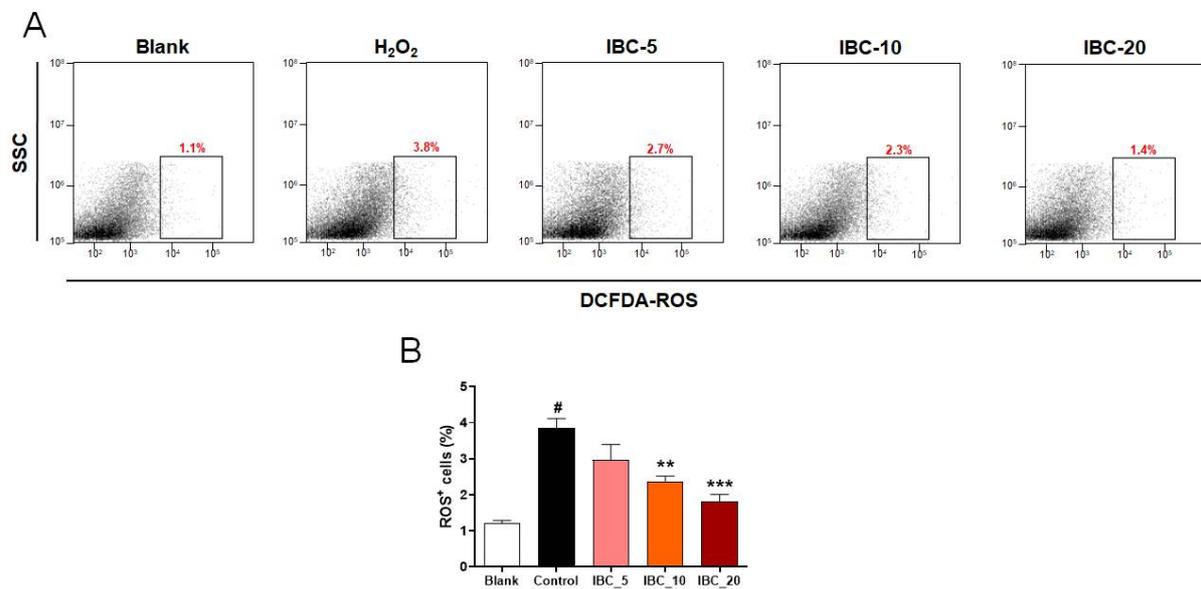


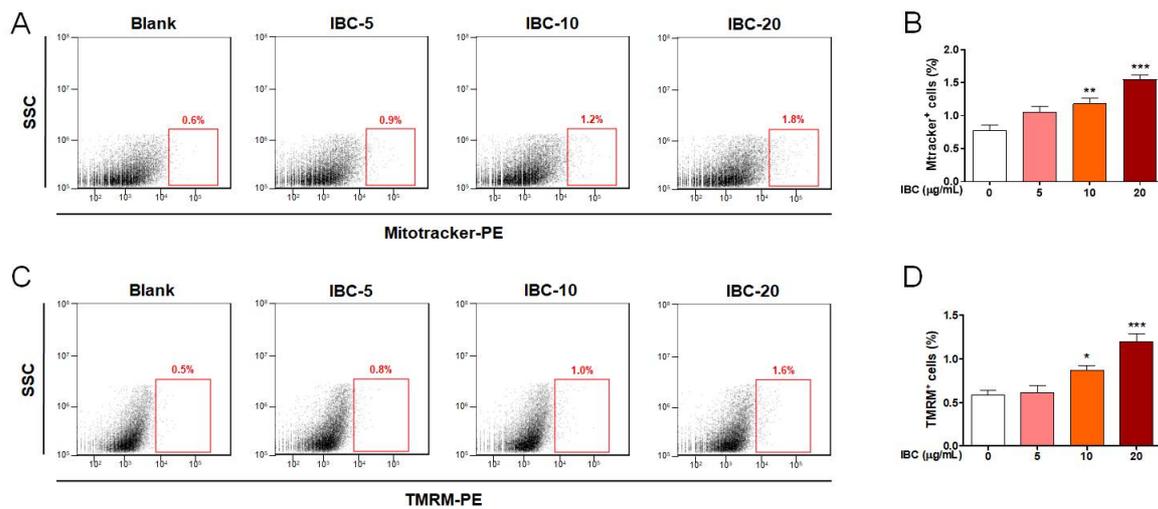
## Supplementary of antioxidants-1096936



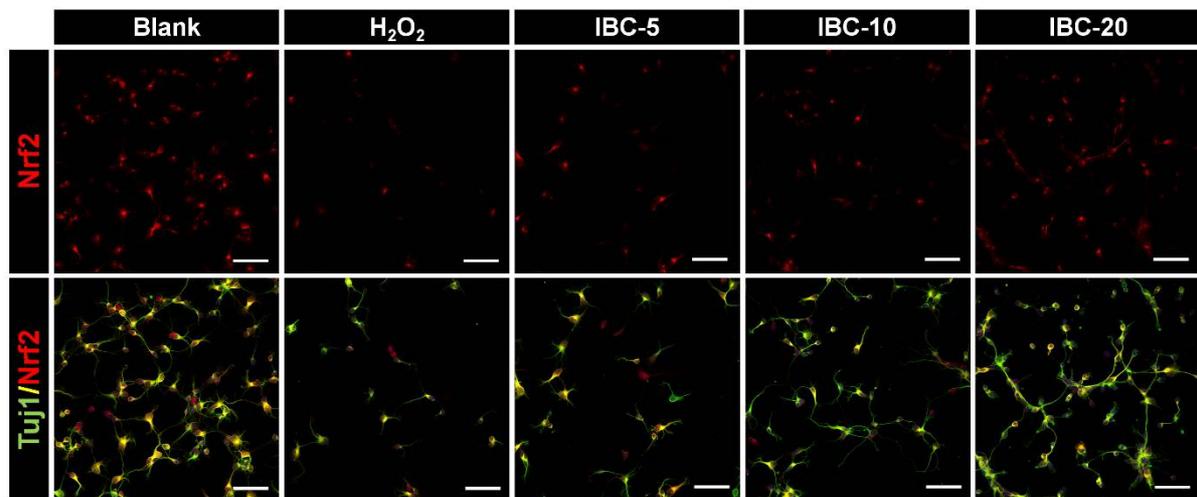
**FigureS1.** Immunocytochemical analysis of cortical neurons to assess the H<sub>2</sub>O<sub>2</sub> optimal concentration and incubation time. Representative immunocytochemical images showing TuJ1 (green) in cortical neurons treated with 100 or 500 μM H<sub>2</sub>O<sub>2</sub> for 30 min or 1 h. White scale bar = 200 μm.



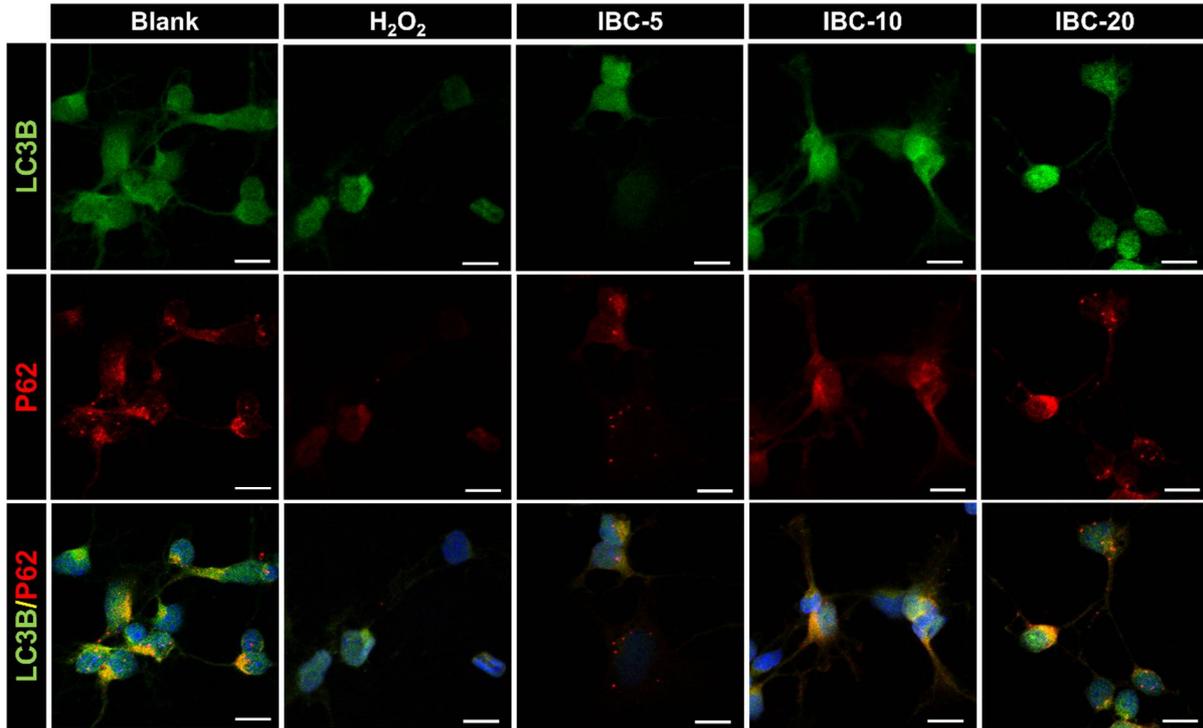
**FigureS2.** Flow cytometric analysis of intracellular ROS in H<sub>2</sub>O<sub>2</sub>-treated cortical neuron with 5, 10 and 20 μg/ml of IBC. (A) Representative flow cytometric dot plots graphs showing DCFDA-ROS expression. (B) Quantification of ROS-positive cells at the single cell level. Data represent mean ± SEM of six independent experiments. Significant differences indicated as #  $p < 0.001$  compared vs. the blank group, \*\*  $p < 0.01$  and \*\*\*  $p < 0.001$  vs. the H<sub>2</sub>O<sub>2</sub> group were analyzed via one-way ANOVA with Tukey's post-hoc test.



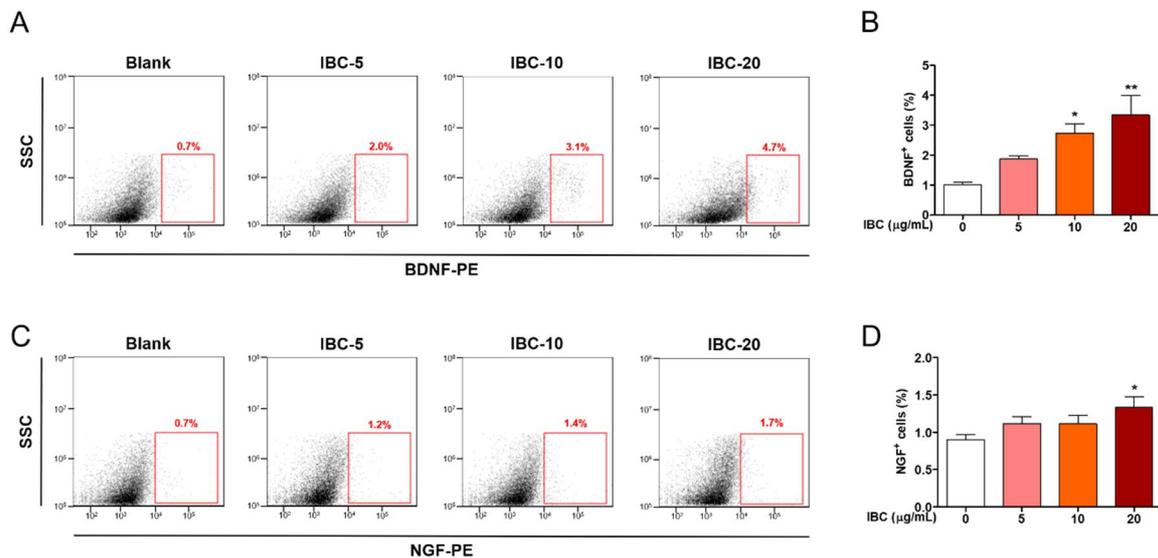
**FigureS3.** Mitotracker (Mtracker) and TMRM Flow cytometry analysis of cortical neuron treated with 5, 10 and 20  $\mu\text{g/ml}$  of IBC without  $\text{H}_2\text{O}_2$  exposure. (A) Representative dot plots graphs showing Mtracker expression. (B) Quantification of Mtracker-positive cells at the single cell level. (C) Representative dot plots graphs showing TMRM expression. (D) Quantification of TMRM-positive cells at the single cell level. Data represent mean  $\pm$  SEM of six independent experiments. Statistically significant differences compared with blank group (0  $\mu\text{g/ml}$ ) by one-way ANOVA with Tukey's post-hoc test. (\* $p < 0.05$ ; \*\* $p < 0.01$ ; \*\*\* $p < 0.001$ ).



**FigureS4.** Nrf2 immunocytochemical analysis of cortical neuron treated with 5, 10 and 20  $\mu\text{g/ml}$  of IBC with  $\text{H}_2\text{O}_2$  exposure. Representative immunocytochemical images showing Nrf2 (red) and Tuj1 (green) in  $\text{H}_2\text{O}_2$ -treated cortical neurons with 5, 10 and 20  $\mu\text{g/ml}$  of IBC. White scale bar = 50  $\mu\text{m}$ .



**FigureS5.** LC3B/P62 immunocytochemical analysis of cortical neurons treated with 5, 10 and 20  $\mu\text{g/ml}$  of IBC with  $\text{H}_2\text{O}_2$  exposure. Representative immunocytochemical images showing LC3B (green) and p62 (red) in  $\text{H}_2\text{O}_2$ -treated cortical neurons with 5, 10 and 20  $\mu\text{g/ml}$  of IBC. White scale bar = 15  $\mu\text{m}$ .



**FigureS6.** BDNF and NGF Flow cytometry analysis of cortical neuron treated with 5, 10 and 20  $\mu\text{g/ml}$  of IBC without  $\text{H}_2\text{O}_2$  exposure. (A) Representative dot plots graphs showing BDNF expression. (B) Quantification of BDNF-positive cells at the single cell level. (C) Representative dot plots graphs showing NGF expression. (D) Quantification of NGF-positive cells at the single cell level. Data represent mean  $\pm$  SEM of six independent experiments. Statistically significant differences compared with blank group (0  $\mu\text{g/ml}$ ) by one-way ANOVA with Tukey's post-hoc test. (\* $p < 0.05$ ; \*\* $p < 0.01$ ).