

랫드대뇌피질 신경세포에 있어서 과산화수소로 유발된 산화적 신경세포손상에 대한
지유의 보호효과

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**Protective effect of Sanguisorbae Radix on H₂O₂-induced oxidative neuronal
cell damage in rat cortical neurons**

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Objectives

Sanguisorbae radix (SR) from Sanguisorba officinalis L. (Rosaceae) is widely used in Korea and China due to its various pharmacological activity. We previously reported that SR prevent A β (25-35)-induced neuronal cell damage *in vitro*. The present study aims to investigate the effect of the methanol extract of SR on H₂O₂-induced neurotoxicity using cultured rat cortical neurons.

Materials and Methods

- Materials
 - Methanol extract of SR
 - H₂O₂
- Methods
 - Primary neuronal culture: Cerebral cortical neurons (E15 SD rat)
 - Analysis of cell viability: MTT colorimetric assay, Hoechst 33342 staining
 - Measurement of Glutamate release: HPLC - ECD

Results and discussions

SR, over a concentration range of 10-50 μ g/ml, inhibited H₂O₂-induced neuronal cell death, as assessed by MTT assay and the number of apoptotic nuclei, evidenced by Hoechst 33342 staining. Pretreatment of SR (10, 30 and 50 μ g/ml) inhibited glutamate release into medium induced by 100 μ M H₂O₂, which was measured by HPLC. In conclusion, we demonstrated that SR prevents H₂O₂-induced neuronal cell damage *in vitro*.

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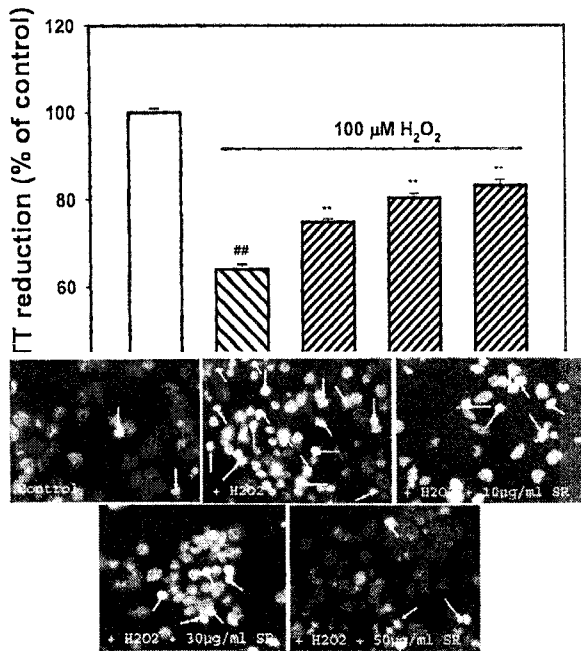


Fig. 1. Inhibitory effect of SR on H₂O₂-induced neuronal cell death in cultured cortical neurons. Neuronal death was measured by the MTT assay. ## p<0.01 compared to control. ** p<0.01 compared to 100 μM H₂O₂.

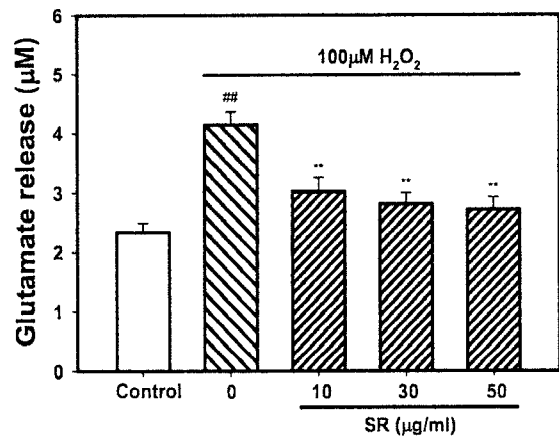


Fig. 2. Inhibitory effect of SR on H₂O₂-induced glutamate release. ## p<0.01 compared to control. ** p<0.01 compared to 100 μM H₂O₂.

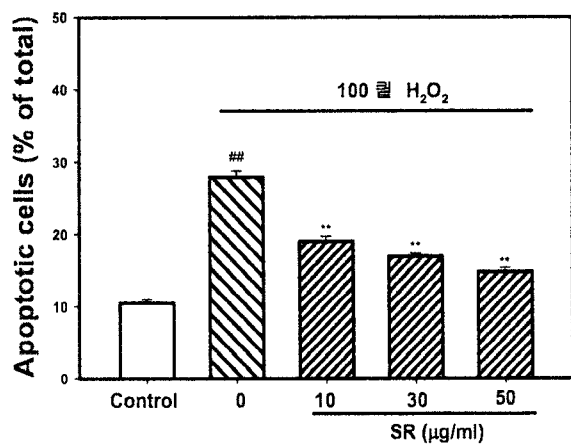


Fig. 3. Inhibitory effect of SR on H₂O₂-induced apoptosis of cultured cortical neurons as measured by Hoechst 33342 staining. ## p<0.01 compared to control. ** p<0.01 compared to 100 μM H₂O₂.