cause to develop aging and various adult diseases. Development of antioxidants to reduce or prevent the diseases associated with free radical have been widely investigated. This studied was carried out to investigate the antioxidative and antihepatic activities of Galla Rhois (*Rhus javanica Linne*). The major components were isolated from the n-Hexane, EtOAc and BuOH extract of Galla Rhois. Their strutures were characterized as syringic acid, gallic acid, gallic acid methyl ester, protocatechuic acid and 1,2,3,4,6-penta-O-galloyl-β-D-glucose by chemical and physical analysis. Five ompounds were tested for antihepatic protective effects on CCl₄ induced cytotoxicity in primary cultured rat hepatocytes and antioxidative effect on DPPH, ferric-thiocyanate method and TBA method.

[PD2-32] [04/21/2000 (Fri) 14:50 - 15:50 / [1st Fl, Bldg 3]]

Curcumin-Induced Inhibition of Microglial Activation

Jung KKO, Yi SW, Kang SY, Kim TG, Kang JH, Kim SH and Han HM

Dept. of Pharmacology, NITR, KFDA

Microglia, brain resident macrophages, play a central role in the inflammatory responses of the brain and are activated in brain injuries and several neurodegerative diseases such as Alzheimer's and Huntington's disease, thereby aggrevating the course of these diseases. In this study, the effect of plant-derived compounds such as curcumin or gingerol on the microglial activation was examined. Microglial cultures were prepared from $2\sim3$ week mixed primary glial cultures obtained from the cerebral cortex of $1\sim2$ day old rats and identified by immunocytochemistry using microglial–specific antibody OX-42. Microglia were activated by lipopolysaccharide(LPS) and interferon– χ (IFN- χ) and the effect of curcumin or 6-gingerol on the microglial activation was examined. Specific parameters measured to monitor microglial activation were nitric oxide(NO) release, prostaglandin E2(PGE2) release and tumor necrosis factor– χ (TNF- χ) release. Curcumin($1\sim10\mu$ M) inhibited NO release induced by LPS and IFN- χ in a dose–dependent manner whreas 6-gingerol($2\sim20\mu$ M) did not have effect on LPS plus IFN- χ -induced NO release. The levels of PGE2 and TNF- χ induced by LPS and IFN- χ were also inhibited by $1\sim10\mu$ M curcumin in a dose–dependent manner. These results showed that curcumin could modulate microglial activation.

[PD2-33] [04/21/2000 (Fri) 14:50 - 15:50 / [1st Fl, Bldg 3]]

Potential Protective Effects of Phytochemicals on the Apoptosis induced by Ultralviolet Irradiation and Oxidative Stress

Kim TGO, Park HJ, Han HM, Kang SY, Jung KK, Kang JH, Lee EA and Kim SH

Dept. of Pharmacology, NITR, KFDA

Exposure to ultraviolet (UV) irradiation and oxidative stress like reactive oxygen species (ROS) induce a celluar injury involving an apoptotic cell death in living organisms. Therefore, the removal of ROS and the inhibition of apoptotic cell death might be a critical step for preventing some pathological conditions including aging and cancer. In this study we examined some phytochemicals to find out whether they had an effect on scavenging of hydroxyl radical or superoxide anion and inhibiting of H2O2- or UV-induced apoptosis. The extract of aloe, Rhus verniciflua (MU2) and Ulmus davidiana showed the protective effects for the hydroxyl radical-induced DNA strand break and the dose-dependent scavenging effects for hydroxyl radical and superoxide anion. Aloin showed the protective effect for DNA strand break and aloesin for hydroxyl radicals, respectively. The effects on the oxidative stress-induced apoptosis in NIH 3T3 fibloblast had been examined. Aloe extract, Aloesin and MU2 showed inhibition for the H2O2-induced apoptosis, which were consistent with the