in previous reports, we exhibited that acteoside showed significant cytotoxicity against various cancer cells. In this study we investigated that acteoside is capable of inducing differentiation in HL-60 human leukemia cell line. After being treated with acteoside, the growth curve was decreased remarkably in a dose- and time-dependent manner, and cell doubling time was delayed. Exposure of cells to 20 μg/ml acteoside induced differentiation of HL-60 cells to monocyte/macrophage-like cells by cell surface antigen expression. The percentage of NBT reducing activity was increased in a time-dependent manner. In addition, the protein level of p21 and p16 increased and pp60 decreased in western blot analysis. These results suggest that acteoside possess the activity of inducing differentiation in HL-60 cells.

[PC1−15] [ 10/17/2002 (Thr) 13:30 – 16:30 / Hall C ]

An investigation of the effect of epigallocatechin-3-gallate on the renal dipeptidase release

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The action of epigallocatechin-3-gallate (EGCG), polyphenol compound from green tea, on the release pattern of glycosylphosphatidylinositol (GPI)-anchored renal dipeptidase (RDPane) from renal proximal tubules (PTs) was examined. EGCG had a stronger inhibitory effect on the release of RDPane than alkaline phosphatase (APase), another GPI-anchored ectoenzyme used as a reference protein. The effect of EGCG on cell viability as assessed by MTT test was found to be intact, and moreover, was indicative of potent cell activation or proliferation. Inhibition of RDPane release by EGCG was also confirmed by time-dependent manner. The concentration of nitric oxide (NO), measured by nitrite, in the incubation supernatant increased up to 6-fold at 100μM EGCG, whereas the release of RDPane decreased in inverse proportion less than 10% of the control, thus demonstrating the inhibitory effect of EGCG on the release of RDPane via NO production. We investigated whether the inhibitory effect of EGCG by NO production was affected by NO-dependent downstream cGMP signaling with the specific inhibitors of the NO-soluble guanylate cyclase pathway, 1H-[1,2,4]oxadiazolo[4,3-a]quinazolin-1-one (ODQ) and the inhibitor of cGMP-selective phosphodiesterase, Zaprinast. However, there was no evidence of cGMP involvement and also of intracellular Ca²⁺ concentration in inhibition of RDPane release by EGCG. When the supernatant PLC by incubation PTs was treated to PTs with EGCG for 15min, the marked increase of RDPane release was strongly blocked. These data support that the inhibitory action of EGCG on the RDPane release was mainly determined by an interference of external Pi-PLC reaction rather than cellular signaling stimulated by NO.

[PC1−16] [ 10/17/2002 (Thr) 13:30 – 16:30 / Hall C ]

Peroxynitrite Scavenging Activity of Active Constituents from Scutellaria baicalensis

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Peroxynitrite, formed from the reaction of .O₂⁻ and .NO, is a cytotoxic species that can oxidize several cellular components such as proteins, lipids and DNA. Oxidative stress is considered to be the major cause of aging and many age-related diseases including Alzheimer's disease, rheumatoid arthritis, cancer, and atherosclerosis. ONOO−, a powerful oxidant, can cause damage of proteins, lipid and DNA through nitration and oxidation. The aim of this study was to evaluate the ability of methanolic extract and fractions from Scutellaria baicalensis and to screen the active components. Methanolic extract showed strong ONOO− scavenging activity. Among fractions, ethylacetate fraction had the potent scavenging activity. In further analysis, baicalin, 5,7,2',5'-tetrahydroxy-8,6'-dimethoxyflavone(TDF), wogon, baikalain, rhamontigenin and rhamonticin were identified from fractionated extract. Results showed that the most effective compounds were baicalin and TDF which led to decreased ONOO− mediated nitration of tyrosine through electron nitration and showed significant inhibition on nitration of albumin and GSH reduction by ONOO− in a dose-dependent manner. TDF and baikalain can be developed as novel ONOO− scavengers for the prevention of ONOO− involved diseases.