

dipyridyl diselenide on iNOS and COX-2 expression induced LPS(lipopolysaccharide) in Raw 264.7 murine macrophages. This organoselenium compounds inhibited NO production, iNOS expression, and COX-2 expression in a concentration-dependent manner. These findings suggest that this organoselenium compounds exert anti-inflammatory effect by inhibiting expression of iNOS and COX-2.

[PC1-6] [04/18/2002 (Thr) 14:00 - 17:00 / Hall E]

Development of Protein Chip for Osteoporosis Diagnosis

Kim So Young^o, Yun Mi Yong, Chang Dong Il, Lee Nam Taek, Choi Myung Ja

Bioanalysis & Biotransformation Research Center, Korea Institute of Science & Technology, Department of Chemistry, Korea Military Academy

Osteoporosis has been characterized by its low bone mass and architectural deterioration of bone tissue. We performed noncompetitive enzyme immunoassay (EIA) for the detection of osteoprotegerin (OPG) and transforming growth factor β 3 (TGF β 3) as osteoporotic biomarkers. Sensitive EIA calibration curves were obtained for OPG and TGF β 3 with detection limits of 0.05 ng/mL and 0.1 ng/mL, respectively, under optimized condition. To develop a solid matrix of protein chip, we initially treated a silica-based glass with 3-aminopropyltriethoxysilane (APS), and additionally with either glutaraldehyde or sulfo-EGS (Ethyleneglycol bis(sulfosuccinimidyl-succinate)) to activate the amino group of APS. The antibody of osteoporotic biomarkers were immobilized on the chip surfaces and calibration curves for OPG and TGF β 3 using a microchip were compared with those obtained from EIA. We found that detection limits performed with microchip considerably correlated with those of EIA. These results promisingly indicate that the microchip can be applied for the diagnostic tool of osteoporosis as EIA.

[PC1-7] [04/18/2002 (Thr) 14:00 - 17:00 / Hall E]

Alpha-viniferin: paw edema reduction and down-regulation of inflammatory mediators

Chung Eun Yong^o, Min Kyung Rak, Kim Youngsoo

College of Pharmacy, Chungbuk National University

Anti-inflammatory activity of alpha-viniferin, a polymeric compound of resveratrol, has been demonstrated in an animal model, and inhibitory effect of the compound on inflammatory mediators has been investigated in order to elucidate mode of the action. When administered orally with >30 mg/kg or injected intravenously with >3 mg/kg, alpha-viniferin showed significant anti-inflammatory activity on carrageenin-induced paw edema in mice. Alpha-viniferin showed IC50 values of 4.9 μ M on cyclooxygenase (COX)-2 activity, 2.7 μ M on production of nitric oxide in lipopolysaccharide (LPS)-stimulated murine macrophages Raw264.7, and 8.5-9.8 μ M on production of superoxide anions in unopsonized zymosan-stimulated human monocytes and neutrophils. The compound showed very weak inhibitory effect on COX-1 and myeloperoxidase activities. Alpha-viniferin showed differential inhibitory effects on proinflammatory cytokines with IC50 values of 10.4 μ M on interleukin (IL)-3 bioactivity, 18.9 μ M on IL-5 bioactivity, and 18.8 μ M on IL-6 bioactivity. Alpha-viniferin showed an IC50 value of 9.8 μ M on tumor necrosis factor (TNF) production in LPS-stimulated Raw264.7 cells, but did not inhibit the IL-1 and TNF bioactivities. These pharmacological findings expand the importance of alpha-viniferin as a beneficial agent to human health, and will help to clarify protective mechanisms of the compound against inflammatory conditions.

[PC1-8] [04/18/2002 (Thr) 14:00 - 17:00 / Hall E]

Antioxidant Effect of Flavonoids and Phenolic Acids on Early Phase of Cu²⁺-Catalyzed LDL Oxidation

Kim JuRyoung^o, Jeong Taesuk, Sok DaiEun

College of Pharmacy, Chungnam National University, Taejon 305-764, Korea *Korea Research Institute of Bioscience and Biotechnology, Taejon 305-333, Korea

There have been increasing evidences that oxidative modification of low density lipoprotein leads to the formation of cholesterol deposits in foam cell, which is related to atherogenesis. Although various antioxidants have been employed to prevent against LDL oxidation, the study concerning the effect of antioxidants on the early phase of LDL oxidation is limited. Here, the effects of flavonoids and phenolic acids, possessing a *o*-dihydroxy moiety, were studied on Cu^{2+} (10 μM)-catalyzed LDL oxidation in early stage (20 min) by measuring the formation of peroxide by chemiluminescence method. The order of antioxidant activity was catechin > quercetin = caffeic acid > luteolin > gallic acid, with catechin (IC_{50} , 0.21 μM) being the most potent. Separately, the antioxidant action was evaluated by measuring the TBA value during 4 hr LDL oxidation. Generally, there was a good correlationship of potency between peroxide value in early stage and TBA value during 4 hr oxidation, the antioxidant activity seemed to be stronger (approximately 5-folds) in early step than late step. Especially, the antioxidant action of some antioxidants differed greatly according to oxidation time. Further studies remain to be performed in order to assess the combinational effect of flavonoids and phenolic acids on the early phase of Cu^{2+} -catalyzed LDL oxidation.

[PC1-9] [04/18/2002 (Thr) 14:00 - 17:00 / Hall E]

Peroxynitrite Scavenging Activity of Sinapic Acid

Zou YaNi^o, Kim AeRa, Kim JungEun, Park TaeHyun, Choi JaeSue, Chung HaeYoung

College of Pharmacy, Pusan National University, Pusan 609-735, Korea, and Food Science and Biotechnology, Pukyong National University, Pusan 608-737, Korea

Peroxynitrite (ONOO^-), formed from a reaction of superoxide (O_2^-) and nitric oxide (NO), is one of most potent cytotoxic species that are known to oxidize cellular constituents including essential proteins, lipids and DNA. In this study, the ability of sinapic acid (3,5-dimethoxy-4-hydroxycinnamic acid), isolated from *Brassica juncea*, to scavenge ONOO^- was investigated. The data obtained show that sinapic acid can efficiently scavenge native ONOO^- as well as ONOO^- derived from peroxynitrite donor 3-morpholinopropanone hydrochloride (SIN-1). Spectrophotometric analyses revealed that sinapic acid suppressed the formation of ONOO^- -mediated tyrosine nitration through electron donation mechanism. In further studies, sinapic acid also showed a significant ability inhibiting nitration of bovine serum albumin (BSA) and low-density lipoprotein (LDL) in a dose-dependent manner. Sinapic acid decreased the LDL peroxidation induced by SIN-1-derived ONOO^- . The present study documented that sinapic acid has an efficient ONOO^- scavenging ability, which may be a potent ONOO^- oxidant scavenger for the protection of the cellular defense activity against the ONOO^- -involved diseases.

[PC1-10] [04/18/2002 (Thr) 14:00 - 17:00 / Hall E]

Effect of Carvacrol, a Terpenoid of Black Cumin Oil on the Arachidonic Acid Metabolism in Rabbit Platelet Aggregation

Son DongJu^o, Im KyungHa, Satoshi Akiba*, Takashi Sato*, Park YoungHyun

College of Natural Sciences, Soonchunhyang University, Asan, Korea * Department of Pathological Biochemistry, Kyoto Pharmaceutical University, Kyoto, Japan

Black cumin, the seeds of *Nigella sativa* L., has been used in Arab countries as food and traditional medicine, which has therapeutic effects on various diseases such as asthma, flatulence, polio, kidney stones and abdominal pain. The immunopotential, anti-tumor, anti-inflammatory, anti-hypertension, hypoglycemia, respiratory stimulation, anti-oxytocic, and anti-bacterial effects were reported. In this study, the effects of carvacrol, a terpenoid of black cumin oil, on platelet aggregation and arachidonic acid (AA) metabolism have been investigated using washed rabbit platelets. AA liberation and generation of thromboxane B_2 (TXB_2), prostaglandin D_2 (PGD_2), and 12-hydroxyeicosatetraenoic acid (12-HETE) were evaluated by radio-chromatographic analysis with washed rabbit platelet in vitro. Carvacrol inhibited