

¹Research Center for Proteineous Materials, Chosun University, Gwangju 501-759, Korea,

¹College of Pharmacy, Chosun University, Gwangju 501-759, Korea

The genus *Styrax* (Styracaceae) is different from other genera of this family due to the production of resinous material, usually secreted when the barks and trunks are injured by sharp objects. This resin, in the past considered a miraculous remedy in several parts of Asia and America, has been used in traditional medicine to treat inflammatory diseases. The CH₂Cl₂ fraction of *Styrax japonica* showed significant cytotoxic activities by SRB method against five human tumor cell lines (A549, HCT-15, MES-SA, SK-OV-3, and SK-MEL-2). We isolated four known pentacyclic triterpenoids by bio-activity guided fractionation and identified as oleanolic aldehyde acetate (1), euphoringinol (2), erythrodiol-3-acetate (3), and anhydrosophoradiol (4). Compounds 1-4 were isolated from *S. japonica* for the first time. The triterpenoids were identified by comparison with spectroscopic data. And we also were assayed for cytotoxic activities of compounds 1-4.

[PD2-52] [2003-10-11 09:00 - 12:30 / Grand Ballroom Pre-function]

Platelet Anti-aggregating Triterpene and Sterol Constituents from the Leaves of *Acanthopanax senticosus*

Jin Jing Ling^o, Lee Sanghyun, Lee Yong Yook, Kim JeongMi, Heo Jung Eun, Yun-Choi Hye Sook

Natural Products Research Institute, Seoul National University, Seoul 110-460, Korea

From methanol extract of *Acanthopanax senticosus*, six platelet anti-aggregating compounds, chiisanogenin (1), chiisanoside (2), ursolic acid (3), oleanolic acid (4), b-sitosterol (5) and daucosterol (6) were isolated. All of the isolated compounds showed dose-dependent inhibitory activities to rat platelet aggregation induced by all the agonist employed. Compound 1 showed about 50 folds higher potency than acetylsalicylic acid (ASA) on U46619 induced platelet aggregation (IC₅₀: 6.21 μM) and 10 ~ 20 folds higher effect than ASA on epinephrine and arachidonic acid (AA) induced aggregation (IC₅₀: 2.50 and 4.81 μM, respectively). Compounds 5 and 6 were 2 ~ 6 folds more inhibitory than ASA on collagen (IC₅₀: 195 and 114 μM respectively) and U46619 (IC₅₀: 170 and 56.1 μM respectively) induced aggregation.

[PD2-53] [2003-10-11 09:00 - 12:30 / Grand Ballroom Pre-function]

Cholinesterase-inhibitory Farnesylacetone Derivatives from the Brown Alga *Sargassum sagamianum*

Park Soo Hee^o, Hwang Jeong Won, Lee Bong Ho, Choi Byoung Wook, Ryu GeonSeek

Dept. of Chemical Technology, Hanbat National University

In continuing search for bioactive compounds from Korean marine algae, we found cholinesterase-inhibitory activity in the methanolic extract of brown alga *Sargassum sagamianum*. After partitioning between CHCl₃ and 30% MeOH, the former layer was purified by a series of ODS flash, silica column, gel-filtration on Sephadex LH-20, and HPLC to give two farnesylacetone derivatives. Their structures were identified by comparison with the literature data. Compounds 1 and 2 showed moderate acetylcholinesterase and butyrylcholinesterase inhibitory activities with IC₅₀ values of 65.0~48.0 μM and 34.0~23.0 μM, respectively. Interestingly, farnesylacetones have different skeletons from the known cholinesterase inhibitors such as tacrine, physostigmine, huperzine A, donepezil and tolserine.

[PD2-54] [2003-10-11 09:00 - 12:30 / Grand Ballroom Pre-function]

Melanin Biosynthesis Inhibitors from the Tubers of *Gastrodia elata*

Li Gao^o, Kim Jae-Hyon, Xu Minglu, Seo Chang-Seob, Kim HyoJin, Lee YouJeong, Lee YeunKoung, Lee Seung-Ho, Chang Hyeun Wook, Son Jong-Keun

College of Pharmacy, Yeungnam University

The bioassay-guided fractionation of the methylene chloride soluble portion of a methanol extract of *Gastrodia elata* tubers led to the isolation of a new furfural, 5-(4-hydroxy-benzyloxymethyl)-furan-2-carbaldehyde (2), together with four known compounds (1, 3-5), which exhibited potent inhibitory activity at the concentration of 25 µg/ml on melanin biosynthesis in cultured B-16 mouse melanoma cells.

[PD2-55] [2003-10-11 09:00 - 12:30 / Grand Ballroom Pre-function]

The antioxidative compounds of the *Aster tataricus*

Choi Doo-Youn¹o, Moon Young-Hee², Woo Eun-Rhan^{1,2}

¹*Research Center for Proteineous Materials, Chosun University, Gwangju 501-759, Korea*

²*College of Pharmacy, Chosun University, Gwangju 501-759, Korea,*

^{1,2}*College of Pharmacy, Chosun University, Gwangju 501-759, Korea*

The *Aster tataricus* is a chinese traditional medicine called "Ziwan" which has an expectorative and remediable cough action. The anti-oxidant activities of *A. tataricus* were investigated. The MeOH extract of *A. tataricus* showed strong anti-oxidant activity in the NBT(nitroblue tetrazolium) method system, and thus fractionated with several solvents in to the EtOAc, n-BuOH, CH₂Cl₂, H₂O fraction. The EtOAc soluble fraction exhibiting strong anti-oxidant activity was further purified by repeated silica gel and sephadex LH-20 column chromatography. Three compounds were isolated from the EtOAc fraction by the activity-oriented purification procedure. Their structures were determined as quercetin, kaempferol, kaempferol 3-O-glucoside, respectively, on the basis of spectral data. The antioxidative compounds of the EtOAc fraction of *A. tataricus* is under study.

[PD3-1] [2003-10-11 09:00 - 12:30 / Grand Ballroom Pre-function]

The compositions of essential oils from *Thymus* species and their antifungal activities

Shin Seungwon^o, Pyun Mi-Sun, Kim Ji-Hyun, Lim Sook, Kim You Sun

College of Pharmacy, Dongduk Women's University

To develop useful antifungal agents from essential oils in Korean plant resources, the activities of *Thymus quinquecostatus* and *T. quinquecostatus* var. *japonica* were evaluated against ten pathogenic fungi. Their results were compared with those of *T. vulgaris*, which is native to Europe. The essential oils of the tested *Thymus* species were obtained by steam distillation using a simultaneous steam distillation-extraction apparatus. The above ground parts of plants cultivated in the herbal garden of Duksung Women's University were used. The composition of the essential oils were analyzed and compared by GC-MS. The antifungal activity of the essential oil fraction of *Thymus* species and thymol, the main component of this oil, were investigated against *Aspergillus niger*, *A. flavus*, *Trichoderma viride*, *Candida albicans*, *C. utilis*, *C. tropicalis*, *Cryptococcus neoformans*, *Trichosporon mucoides*, *Trychophyton tonsurans*, and *Blastoschizomyces capitatus*. The MICs and the growth inhibition against the fungi was evaluated by broth dilution method and disk diffusion test. Additionally, the combination effects of the essential oils with synthetic antibiotics were estimated.

[PD3-2] [2003-10-11 09:00 - 12:30 / Grand Ballroom Pre-function]

Inhibitory effects of Saiko-ka-Ryukotsu-Borei-To on the migration and proliferation of vascular smooth muscle cell and suppression of carotid intimal thickness after balloon injury in rats

Chung Hwa-jin^o, Maruyama Ikuro, Tani Tadato, Lee Sang Kook

Institute of Natural Medicine, Toyama Medical and Pharmaceutical University, Japan, College of Pharmacy, Ewha Womans University, School of Medicine, Kagoshima University, Japan, Institute of Natural Medicine, Toyama Medical and Pharmaceutical University, Japan

Objectives: We have reported that oral administration of Saiko-ka-Ryukotsu-Borei-To (SRB), a traditional