

determined by the interpretation of spectroscopic data obtained from various NMR techniques. A variety of bioassay for them are in progress.

[PD2-5] [04/19/2002 (Fri) 10:00 - 13:00 / Hall E]

A New Uracil Derivative and A New Acylglycosyl Sterol from Quisqualis Fructus

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Quisqualis Fructus is a Korean traditional medicine to treat ascariasis.¹⁾ Quisqualic acid and trigonelline were reported from *Quisqualis Fructus*.²⁾ In the course of our searching for topoisomerase I inhibitor from Korean traditional medicine, *Quisqualis Fructus* exhibited moderate activity. The research of this source led to isolate a new uracil derivative and a new acylglycosyl sterol together with five known compounds. On the basis of spectroscopic data, their structures have been elucidated as 1-(2-amino-1,4-dioxan-3-yl)-uracil, 3-O-[6-O-(8-octadecenoyl)-glycosyl] epicodisterol, 3-amino-acrylamide, pyridylglycine, epicodisterol, betulinic acid and ursolic acid methyl ester. The topoisomerase I inhibitory effect of these compounds are under study.

1) Shanghai Science and Technologic Publisher and Shougakukan, The Dictionary of Chinese Drugs, Shougakukan, Tokyo, pp.1035-1037 (1985)

2) Takemoto, T., Takagi, N. Nakajima, T. and Koike, K., *Yakugaku Zasshi*, 95 (2), 176-179 (1975)

[PD2-6] [04/19/2002 (Fri) 10:00 - 13:00 / Hall E]

Antioxidative activity of *Acanthopanax chiisanensis* Fructus

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Acanthopanax spp. are one of the traditional tonic agents. They have been used as analgesics, stimulant of immune system and replenishment of body functions. In order to estimate the antioxidative activity of *Acanthopanax chiisanensis* Fructus, we measured anti-lipid peroxidative efficacy on human low density lipoprotein (LDL) with TBARS (2-thiobarbituric acid) assay from its MeOH extract. And we tested superoxide scavenging activity by Free radical scavenging assay. Superoxide radicals are generated in a phenazine methosulfate (PMS)-beta-nicotinamide adenine dinucleotide (reduced form, NADH) system by oxidation of NADH and assayed by the reduction of nitroblue tetrazolium (NBT).

[PD2-7] [04/19/2002 (Fri) 10:00 - 13:00 / Hall E]

Antiinflammatory and Antinociceptive Principles of the *Acanthopanax senticosus* Stem Bark

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MeOH extract of the stem bark of *Acanthopanax senticosus* (Araliaceae) was fractionated to test anti-inflammatory in the rat induced by carrageenan and Freund's complete adjuvant reagent (FCA),

respectively. EtOAc-soluble fraction showed the significant inhibition of edema in the two assays. Phytochemical isolation afforded kalopanaxaponin A, hederagenin 3-O-β-D-glucopyranosyl (1-2)-α-L-arabinopyranoside, caffeic acid and liriiodendrin. The three main components, caffeic acid, kalopanaxaponin A and liriiodendrin exhibited significant antiinflammatory action by intraperitoneal administration at 10 mg/kg dose (p<0.01), respectively. The three components also exhibited antinociceptive actions in writhing- and hot plate tests. These results suggest that the stem bark of *A. senticosus* will be applicable for the treatment of rheumatoid arthritis.

[PD2-8] [04/19/2002 (Fri) 10:00 - 13:00 / Hall E]

Phenolic Compounds from *Sophora japonica* wood

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Sophora japonica is called as Chinese scholar tree or Japanese pagoda tree. This species which belongs to Leguminosae family, is used in the traditional medicine. To isolate compounds, column chromatography was used with various solvent system in silica gel and Sephadex LH-20. To identify compounds, instrumental analyses (NMR spectrometry including 1H-1H COSY, NOESY, HMQC, HMBC and Mass spectrometry) were performed.

From the Wood of *S. japonica*, eight phenolic compounds were isolated and identified as follows : irisolidone, biochanin A, formononetin, 7-hydroxy-4'-methoxyisoflavanone, puerol A, 5-hydroxypseudobaptigen-2'-O-β-D-glucopyranoside, biochanin A-7-O-β-D-xylopyranosyl-(1-6)-β-D-glucopyranoside and (-)maackiain. Among these compounds, 5-hydroxypseudobaptigen-2'-O-β-D-glucopyranoside is the new compound.

[PD2-9] [04/19/2002 (Fri) 10:00 - 13:00 / Hall E]

Flavonoid glycosides and coumarins from *Euodia danielli*

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Euodia daniellii HEMSLEY (Rutaceae) is a plant endemic to Korea. It has been used as a folkmedicine for gastric inflammation, extermination of noxious insects, and headache. Four flavonoid glycosides, vitexin, hesperidin and evodioside B from leaves and flavaprin and evodioside B from fruits were isolated. And also, three coumarins, bergapten, xanthotoxin and isopimpinellin from fruits were isolated. Among them, six compounds with the exception of bergapten were isolated from this plant for the first time. Bergapten showed cyclooxygenase-2 inhibitory activity with an IC50 value of 6.2 μg/ml. All the isolates exhibited no cytotoxicity against the human tumor cell lines, A549, SKOV-3, SKMEL-2, XF498, and HCT15.

[PD2-10] [04/19/2002 (Fri) 10:00 - 13:00 / Hall E]

Terpenoid constituents from *Youngia x. koidzumiana*

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Youngia x. koidzumiana is an indigenous plant growing in Mt. Chiri. In our ongoing research for indigenous plant growing in Korea, we investigated the chemical constituents from the MeOH extract of *Y. x. koidzumiana* whole plants. The MeOH extract was partitioned with hexane, ethylacetate, BuOH, successively. Four known compounds (YK-4-C, YK-6-D, YK-10-B, YK-18-B) were isolated from