the folkloric society of Korea, we investigated the isolation of hydrophilic constituents. Chlorogenic acid (1), 3,4-di-O-caffeoylquinic acid (2), 5-O-[1-butyl]-3,4-di-O-caffeoylquinic acid (3) were isolated by combination of silica gel- and ODS column chromatography. The structure of 3 was determined by 1H-1H-, 1H-13C COSY, HMBC and FAB-MS spectra. Compound 3 has not been isolated before from a natural source.

[PD2-15] [10/20/2000 (Fri) 11:30 - 12:30 / [Hall B]]

Phytochemical Constituents of Actinidia arguta

Moon HI, Whang JI, Oh JS, Choi B, Hong SI, Zee OP

Pharmacognosy Lab, College of Pharmacy, SungKyunKwan University

The root of Actinidia arguta was extracted with methanol and the methanol extract was suspended in H2O and successively partitioned with n- Hexane, CH2Cl2, EtOAc and n-BuOH. The repeated column chromatographic separation of the EtOAc extract resulted in the isolation of two flavonoids (compound 2 and 3) and two triterpenes (compound 1 and 4) and CH2Cl2 extract to afford three lignans (compound 5-7). Their structures have been established by spectroscopic methods

[PD2-16] [10/20/2000 (Fri) 11:30 - 12:30 / [Hall B]]

Antigenotoxic and Antimutagenic Activities of a New Component from the Starfish Asterina pectinifera

Lee NJO, Lee DU, Park CH1, Shin YH1, Ham JH2, Han YH2

Department of Biochemistry, Dongguk University, Kyongju 780-714, ¹Department of Pharmacy, Kyungsung University, Pusan 608-736, ²Department of Biology, Dongguk University, Kyongju 780-714, Korea

>From the butanol fraction of the starfish Asterina pectinifera Muller et Troschel (Asteriidae), we have isolated a new component. 5α -cholest-7-en-

3β-ol. Its antigenotoxic and antimutagenic activities were examined by the SOS chromotest with Escherichia coli PQ37 and by Ames test with Salmonella typhimurium TA1538, respectively. 5α-Cholest-7-en-3β-ol showed potent antigenotoxic activity against the mutagens, both MNNG(N-methyl-N'-nitro-N- nitrosoguanidine) and NQO(4-nitroquinoline N-oxide. For 100% of antigenotoxicity, the concentration of the compound applied against MNNG and NQO were 10μg and 5μg per reaction tube, respectively. Its antimutagenic activity with S. typhimurium TA1538 against the mutagen MNNG was very effective. When its concentrations were varied from 1 μg up to 10μg dose per plate, the inhibition ratio of revertant CFU(colony forming unit) of

TA1538 per plate was increased accordingly, from 25.2% to 99.2%. These results suggest that 5 α -cholest-7-en-3 β -ol possesses antigenotoxic and antimutagenic activity and might be useful as a chemopreventive agent.

[PD2-17] [10/20/2000 (Fri) 11:30 - 12:30 / [Hall B]]

Phytochemical investigation of node of the lotus rhizome (Nelumbo nucifera Gaertn.)

Jun-Sik Kim^o, Su-Min Cho, Jee-Hun Kim, Eui-Chan Jung and Min-Won Lee

College of Pharmacy, Chung Ang University, Seoul 156-756, Korea