

Therapeutic Efficacy of YCY on CCl₄-induced Liver Dysfunction in Rats

Hee-Youl Chai, Woon Kwon, Young Min Cho, Ehn Kyoung Choi, Iksoo Kim¹, Kang Sun Ryu¹, Seock-Yeon Hwang², Chi-Young Yun³, Yun-Bae Kim and Jong-Koo Kang

College of Vet. Med., Chungbuk National Univ., Cheongju 361-763, Korea

¹Department of Sericulture and Entomology, NIAST, Suwon 441-857, Korea

²Dept. of Clinical Pathol., Chungbuk National Univ. Hospital, Cheongju 361-711, Korea

³Department of Biology, Daejeon University, Daejeon 300-716, Korea

E-mail: jkkang@cbu.ac.kr

Introduction

Carbon tetrachloride (CCl₄) is known to cause liver injury characterized by centrilobular necrosis. The toxicity is thought to be exerted via cytochrome P-450-mediated metabolism of CCl₄ to trichloromethyl and trichloromethylperoxy radicals, which initiates lipid peroxidation, leading to hepatocellular membrane damage [1, 2]. Our study demonstrates a therapeutic efficacy of YCY, extract of a cricket, *Gryllus bimaculatus*, on liver injuries induced by CCl₄.

Materials and Methods

To examine therapeutic efficacy, liver damage was induced by repeated oral administration of CCl₄ with an initial dose of 1 ml/kg, followed by 0.5 ml/kg twice a week for 10 weeks. The animals were coadministered orally with YCY from 7 weeks during CCl₄ administration, and sacrificed 24 hr after the final administration.

Results and Discussion

The YCY treatment remarkably reduced the increases in liver weights and serum alanine aminotransferase (ALT) and aspartate aminotransferase (AST) levels induced by CCl₄ exposure (Table 1), and attenuated hepatocytic degeneration and fibrosis (data not shown).

Table 1. Effect of YCY on CCl₄-induced liver injuries

Treatment (mg/kg)	Liver weight	ALT	AST
Control(vehicle)	2.82±0.27	59.8±4.3	110.3±14.6
CCl ₄ alone	3.12±0.33	399.5±245.6	596.8±371.8
CCl ₄ +YCY(100)	2.84±0.19	305.3±162.8	546.7±156.2
CCl ₄ +YCY(300)	3.02±0.44	236.0±66.9	320.6±163.3

References

1. Klaassen, C. D. Toxicology. p.887. McGraw-Hill, New York. 2001.
2. Lee, K. Y. et al. J. Appl. Pharmacol. 2002. 10, 12-18.