

Antioxidant and anti-fibrotic properties of *Lythrum salicaria* root in CCL₄-induced liver fibrosis rat model

Seung-Eun Lee^{1†}, Tae-Jin Ahn¹, Young-Ock Kim¹, Jin-Sook Soe¹, Chung-Berm Park¹, Sun-Woo Cha¹, Hae-Young Chung², Nak-Sul Seong¹

¹Department of Herball Crop Research, RDA, Korea

²Department of Pharmacy, Pusan National University, Korea

Objectives

The study was conducted to investigate antioxidant and liver fibrosis protecting activities of *Lythrum salicaria* root

Materials and Methods

- Materials : Root of *L. salicaria* (LSR) was collected in the medicinal crop farm of RDA in 2004.
- Methods
 - Preparation of extract was conducted by extraction procedure with 50% ethanol at 85°C, for 2 hours for the study. Evaporation of the solvent in extracts filtered was conducted under vacuum atmosphere.
 - In vitro antioxidant activity against ROS and peroxy-nitrite in chemical environment and on cells such as YPEN1 and BV2 were analyzed.
 - Animal and treatment : SD male rats were composed of normal group, CCL₄ single treated group (negative control), CCL₄ plus LSR-treated groups (0.5, 0.25g, 0.5g of LSR extract/Kg), CCL₄ plus silymarin-treated groups (0.5, 0.25g, 0.5g of LSR extract/Kg) and induced liver fibrosis with carbon tetrachloride three times for 6 weeks. Liver fibrosis of rats was induced by intra-peritoneal injection of 40% CCL₄ (1.0ml/kg body weight) dissolved in 0.5ml of corn oil twice per week, TBARS production and ratio of GSH/GSSG (reduced glutathione/oxidized glutathione) as antioxidant parameters, hydroxyproline content and microscopic analysis as collagen production indicators were investigated.

Results

Fifty percent ethanol extracts of *L. salicaria* root showed effective *in vitro* antioxidant activities on ROS and peroxy-nitrite. Ratio of hepatic GSH/GSSG in CCL₄ plus LSR extract (0.125~0.5g/Kg)-treated rats increased from 2.8 to 5.7 fold values compared to that of CCL₄-treated rats. Amounts of hydroxyproline in CCL₄ plus LSR extract-administrated rat livers were 5~10µg/mg which were

[†] Corresponding author : Seung-Eun Lee lse1003@rda.go.kr Tel : 043-871-5586

correspond to -44~-72% of the value in CCL₄-treated rat livers (18μg/mg tissue). This collagen reducing effect of liver tissue in CCL₄ plus LSR-treated rats was supported by immunohistochemical and histological observation. The results showed that the root of *L. salicaria* have efficient antioxidant potential and effective antifibrotic activities in liver fibrosis induced rat.

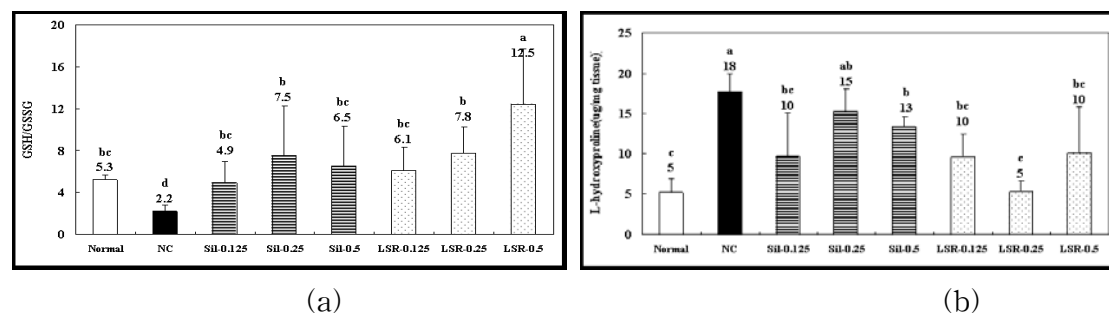


Fig. 1. Effect of LSR extract on the ratio of GSH and GSSH (a) and total collagen content (b) of fibrosis-induced liver of rat

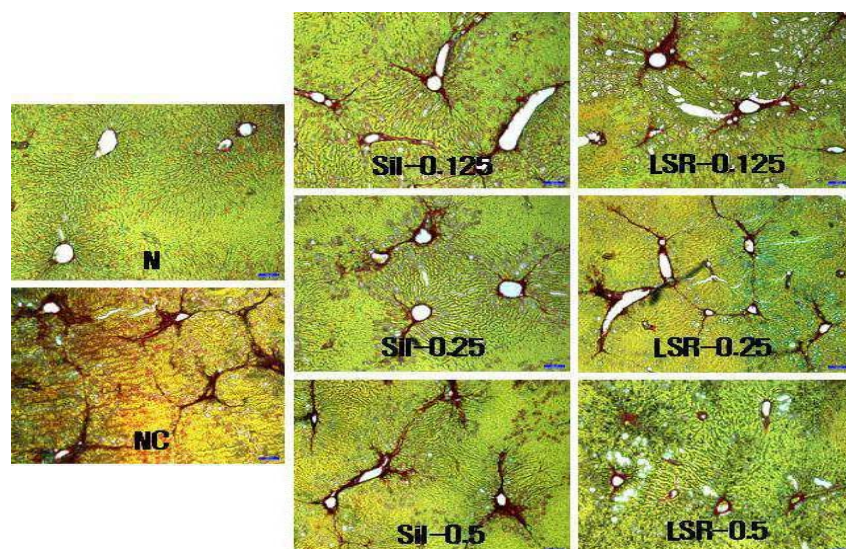


Fig. 2. Histological observation for explaining the effect of LSR extract on collagen production in fibrosis-induced liver of rat (×100). N, vehicle-injected normal group; NC, negative control, CCL₄ (1ml of 40% CCL₄ /Kg body weight with 1.5ml of corn oil plus 1ml of distilled water, 2 times/week for 6 weeks) injected group Sil-0.25, Sil-0.5, 0.25g, 0.5g silymarin/Kg body weight/rat/day-supplemented & CCL₄-injected group; LSR-0.25, LSR-0.5, 0.25g, 0.5g of LSR extract/Kg body weight of rat/day-supplemented & CCL₄-injected group. Values with alphabet on each bar are significantly different at P<0.05.