Hepatotoxicity and nephrotoxicity of saponin-enriched extract of *Asparagus cochinchinensis* in ICR mice

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The inhibitory effects of *Asparagus cochinchinensis* against inflammatory response induced by lipopolysaccharide (LPS), substance P and phthalic anhydride (PA) treatment were recently reported for some cell lines and animal models. To evaluate the hepatotoxicity and nephrotoxicity of *A. cochinchinensis* toward the livers and kidneys of ICR mice, alterations in related markers including body weight, organ weight, urine composition, liver pathology and kidney pathology were analyzed in male and female ICR mice after oral administration of 150, 300 and 600 mg/kg body weight/day saponin-enriched extract of *A. cochinchinensis* (SEAC) for 14 days. The saponin, total flavonoid and total phenol levels were found to be 57.2, 88.5 and 102.1 mg/g in SEAC, respectively, and the scavenging activity of SEAC gradually increased in a dose-dependent manner. Moreover, body and organ weight, clinical phenotypes, urine parameters and mice mortality did not differ between the vehicle and SEAC treated group. Furthermore, no significant alterations were measured in alkaline phosphatase (ALP), alanine aminotransferase (ALT), aspartate aminotransferase (AST), lactate dehydrogenase (LDH), blood urea nitrogen (BUN) and the serum creatinine (Cr) in the SEAC treated group relative to the vehicle treated group. Moreover, the specific pathological features induced by most toxic compounds were not observed upon liver and kidney histological analysis. Overall, the results of the present study suggest that SEAC does not induce any specific toxicity in the livers and kidneys of male and female ICR mice at doses of 600 mg/kg body weight/day.

**Keywords:** *Asparagus cochinchinensis*, hepatotoxicity, nephrotoxicity, saponin, histopathology

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*A. cochinchinensis* is a perennial herb belonging to the Liliaceae family that is widely distributed in China, Japan and Korea, and has long been considered a therapeutic drug because of its anti-inflammatory, diuretic, antiseptic, antitussive, antibacterial, nerve, sialogogue, antipyretic and stomachic effects. *A. cochinchinensis* is also administered in combination with other herbs as a medicine to treat lung disease, immune system-associated diseases and aging [1,2].

Among various therapeutic activities, the anti-inflammatory activity of the extract of *A. cochinchinensis* roots had been investigated in previous studies to generate scientific evidence and found to significantly inhibit production of pro-inflammatory cytokines and tumor necrosis factor (TNF)-α in lipopolysaccharide (LPS)- and substance P-stimulated mouse astrocytes [3].
Moreover, the nitric oxide (NO) concentration decreased in response to compounds 2, 3 and 4 from the ethanol extract of *A. cochinchinensis* roots in LPS-stimulated BV-2 microglial cells. These compounds also greatly decreased indicators of skin inflammation progression including the degree of ectopic edema, ear thickness, cytokine secretion and myeloperoxidase (MPO) activity in a skin inflammation model induced with 12-O-tetradecanoyl-phorbol-13-acetate (TPA) [4]. Furthermore, the aqueous extract of *A. cochinchinensis* suppressed TNF-α-induced cytotoxicity of HepG2 cells [5], enhanced superoxide dismutase (SOD) activity and the spleen index, and decreased the level of malondialdehyde (MDA) in mice [1]. Moreover, saponin-enriched extract of *A. cochinchinensis* (SEAC, ethyl acetate extract) successfully suppressed skin inflammation of IL-4/Luc/CNS-1 transgenic mice induced by phthalic anhydride (PA) treatment [6]. Overall, the results of the aforementioned studies suggest that SEAC can be considered a potential drug for inflammation related diseases in humans. However, its toxicity toward specific organs of mice has not been investigated to date, despite the increased attention and publications regarding the therapeutic effects of SEAC.

Materials and Methods

Preparation of SEAC

The roots of *A. cochinchinensis* used in this study were obtained from Gochang National Agricultural Cooperation Federation in Korea and dried in a drying machine (FD5510S and FD5520S, Ilshinbiobase Co., Dongducheon, Korea) at 60°C. Voucher specimens of *A. cochinchinensis* roots (WPC-14-003) were deposited in the functional materials bank of the Wellbeing RIS Center at Pusan National University. Dried roots were reduced to powder using a pulverizer (MF-3100S, Hanil Electric Co., Seoul, Korea), after which SEAC was obtained at 50°C for 24 h in a 1:10 fixed liquor ratio of solid powder of *A. cochinchinensis* to ethyl acetate solvent ratio using circulating extraction equipment (SHWB-30/45, Woori Science Instrument Co., Pocheon, Korea). These solutions were subsequently passed through a 0.4 μm filter, after which they were concentrated by vacuum evaporation and lyophilization using circulating extraction equipment (IKA Labortechnik, Staufen, Germany) (Figure 1A). Finally, the collected SEAC powder was dissolved in 0.5% Tween 20 Solution (in dH2O) to 600 mg/kg, then further diluted to the required concentration.

Determination of total saponins, phenolic compounds and flavonoids

Total phenolic contents in SEAC were determined using the Folin-Ciocalteu method as previously described [7]. Briefly, a mixture of SEAC solution (1 mL) and Folin-Ciocalteu reagent (5 mL, Sigma-Aldrich Co., St. Louis, MO, USA) was incubated at room temperature for 5 min. This mixture was subsequently added to 15 mL of 20% Na2CO3 and vortexed for 30 sec, after which the absorbance was repeatedly measured at 765 nm using a Versa-max plate reader (Molecular Devices, Sunnyvale, CA, USA). A standard calibration curve was then made using different concentrations of gallic acid (Sigma-Aldrich Co.), and the concentration of total phenolic contents in SEAC was presented as gallic acid equivalent (mg) of extract.

The flavonoid contents in SEAC were measured as previously described [8]. Briefly, 200 μL of several different concentrations of SEAC were mixed with 60 μL of 5% NaNO2 (Sigma-Aldrich Co.) and 60 μL of 10% AlCl3 (Sigma-Aldrich Co.). Following incubation at 25°C, the absorbance was measured using a Versa-max plate reader (Molecular Devices). A standard calibration curve was then made using different concentrations of catechin (Sigma-Aldrich Co.). The flavonoid contents in SEAC are presented as catechin equivalents (mg) of extract.

Finally, the total saponins content was determined using the Helaly method as previously described [9]. Briefly, extract of n-butanol was redissolved in 0.5 mL of 80% MeOH, then mixed with 0.5 mL of 8% vanillin in ethanol and 5 mL of 72% H2SO4 in water. Next, these mixtures were placed in a 60°C water bath for 20 min, then cooled at 0°C for 5 min, after which the absorbance was measured at 544 nm using a Versa-max plate reader (Molecular Devices). The saponin content was calculated from a calibration curve constructed using purified saponin standard (Sigma-Aldrich Co.).
Free radical scavenging activity

The scavenging activity of 2,2-diphenyl-1-picrylhydrazyl (DPPH) radical was measured as previously described [10]. Briefly, each 100 μL sample in eleven different concentrations of SEAC (0-2,000 μg/mL) was mixed with 100 μL DPPH (0.1 mM; Sigma-Aldrich Co.) in 95% ethanol solution or 100 μL 95% ethanol solution, then incubated for 30 min at room temperature. The absorbance of the reaction mixture was subsequently measured at 517 nm using a VersaMax plate reader (Molecular Devices). The DPPH radical scavenging activity of the SEAC was expressed as the percent decrease in absorbance relative to the control. The half maximal inhibitory concentration (IC$_{50}$) was defined as the concentration of substrate that causes a 50% loss in DPPH activity.

Design of animal experiment

The animal protocol used for this experiment was reviewed and approved by the Pusan National University-Institutional Animal Care and Use Committee (PNU-IACUC; Approval Number PNU-2015-0779). All mice were handled in the Pusan National University-Laboratory Animal Resources Center, which is accredited by the Korea Food and Drug Administration (KFDA, Accredited Unit Number-000231) and AAALAC International (Accredited Unit Number; 001525). Seven-week-old female and male ICR mice were purchased from Samtako Bio Korea (Osan, Korea) and provided with ad libitum access to a standard irradiated chow diet (Samtako Bio Korea) consisting of moisture (12.5%), crude protein (25.43%), crude fat (6.06%), crude fiber (3.9%), crude ash (5.31%), calcium (1.14%) and phosphorus (0.99%) and water throughout the feeding study. Before starting the animal experiment, the mice were given at least 1 week to adapt to the experimental environment. During the experiment, mice were maintained in a specific pathogen-free state under a strict light cycle (lights on at 08:00 hr and off at 20:00 hr) at 23±2°C and 50±10% relative humidity.

ICR mice (n=40) were assigned to one of the following four groups: vehicle treated group (n=10), low dosage SEAC treated group (SEACL, n=10), middle dosage SEAC treated group (SEACM, n=10), high dosage SEAC treated group (SEACH, n=10). As a control, one group of ICR mice received an equivalent volume of water via gavage (vehicle treated group), whereas the other groups received 150 mg (SEACL), 300 mg (SEACM) and 600 mg (SEACH)/kg body weight/day of SEAC via oral gavage. The administration period for the toxicity test was based on the Guideline for Drug Toxicity published by the Korea Food and Drug Administration. After SEAC treatment for 14 days, all mice were immediately sacrificed using CO$_2$ gas, after which urine, serum and tissue samples were prepared.

Measurement of body weight and organ weights, and observation of pathological signs

Several pathological signs including dead and abnormal behaviors were recorded more than twice a day for 14 days. In addition, alterations in body weight were observed using an electronic balance (Mettler Toledo, Greifensee, Switzerland) every day according to the KFDA guidelines. The weights of eight organs (brain, ovary (or testis), kidney, spleen, liver, thymus, heart and lung) collected from the sacrificed mice were determined using the same method employed to detect the body weight.

Urine analysis

All mice were sacrificed using CO$_2$ gas at 24 h after the final administration, at which time urine was collected from their bladders and assayed for bilirubin, urobilinogen, ketones, protein, pH, specific gravity and leucocytes using an URiSCAN optima II urine analyzer (Yeongdong Electronics Co., Ltd., Yongin, Korea). All assays were conducted in duplicate using fresh urine.

Serum biochemical analysis

After fasting for 8 h, whole blood from each mouse in all groups was collected from their abdominal veins and incubated for 30 min at room temperature in a serum separating tube (BD Container, Franklin Lakes, NJ, USA). Serum was then obtained by centrifugation at 1,500×g and analyzed for alkaline phosphatase (ALP), alanine aminotransferase (ALT), aspartate aminotransferase (AST), lactate dehydrogenase (LDH), blood urea nitrogen (BUN) and creatinine (Cr) using an automatic biochemical analyzer (BS-120, Mindray, China). All assays were conducted in duplicate using fresh serum.

Histological analysis

Liver and kidney tissues were dissected from mice and fixed in 4% neutral buffered formaldehyde (pH 6.8) overnight, after which each liver was dehydrated and embedded in paraffin wax. Next, a series of liver sections (4 μm) was cut from paraffin-embedded tissue using a
Leica microtome (Leica Microsystems, Wetzlar, Germany). These sections were then deparaffinized with xylene, rehydrated with ethanol at a graded decreasing concentration of 100-70%, and finally washed with distilled water. The slides with liver sections were stained with hematoxylin (Sigma-Aldrich Co.) and eosin (Sigma-Aldrich Co.), then washed with dH₂O, after which pathological changes were measured using the Leica Application Suite (Leica Microsystems).

Statistical analysis
Significant differences between vehicle and SEAC treated ICR mice were identified by one-way analysis of variance (ANOVA) using SPSS for Windows, Release 10.10, Standard Version (SPSS Inc., Chicago, IL, USA). All values are reported as the means±standard deviation (SD) and a p<0.05 was considered significant.

Results

Biochemical properties of SEAC
The main components of SEAC were analyzed to verify their biochemical properties. Among these compounds, crude saponins was found to be present in the greatest abundance (57.2 mg/g), while the levels of flavonoid and polyphenol were 88.5 and 102.1 μg/g, respectively (Figure 1). To investigate the inhibitory activity of SEAC against oxidative stress, the scavenging activity of SEAC against DPPH radicals was measured. The scavenging activity was gradually increased in a dose-dependent manner and the IC₅₀ value was 835.90 μg/mL (Figure 1C). Therefore, the present results suggest that SEAC contain various compounds with high antioxidant activity.

Clinical phenotypes and mortality of SEAC treated ICR mice
To investigate whether SEAC administration affects phenotypic toxicity, the clinical phenotypes and mortality were observed in male and female ICR mice for 14 days. No mice treated with any of the investigated concentrations of SEAC showed significant pathological symptoms including body weight loss, hypokinesia, melancholy, gait abnormality and tremors. No significant differences in body weight were observed between the vehicle and SEAC treated group throughout the 14 day experimental period, although the male group was higher than female group (Table 1). Moreover, dead mice were not observed in the SEAC treated group throughout the experimental period. These results show that SEAC does not induce any significant pathological symptoms or mortality and thus the minimum lethal dose (MLD) is higher than 600 mg/kg of SEAC.

Alteration of organ weights and urine parameters in SEAC treated ICR mice
To investigate the effects of SEAC administration on the weights of organs and urine composition of animals, the weights of eight organs and seven urine factors were measured in male and female ICR mice treated with SEAC. No significant differences in the weights of any of the organs were observed between the vehicle and SEAC treated groups (Table 1). Moreover, the SEAC treated ICR mice showed no morphological changes in the brain, ovary (or testis), kidney, spleen, liver, thymus, heart and lungs when compared to the vehicle treated group. Finally, similar levels of urine parameters (bilirubin, urobilinogen, leucocytes, protein, pH, specific gravity and ketones) were measured in the vehicle and SEAC treated group (Table 2). These results suggest that SEAC treatment for 14 days does not induce any toxicity toward organ weights or urine parameters in male and female ICR mice.

Effects of SEAC on hepatotoxicity of male and female ICR mice
To investigate whether SEAC treatment for 14 days induced hepatotoxicity in ICR mice, alteration of several enzymes related to liver metabolism were measured in the serum of male and female ICR mice treated with SEAC. No significant differences in liver toxicity indicators such as ALP, AST, ALP and LDH were observed between the vehicle and SEAC treated group (Table 3). Furthermore, we investigated whether the lack of alterations of toxicity indicators in serum could completely reflect the histopathological structure of liver tissue. To achieve this, liver sections were stained with H&E and observed microscopically. There were no significant pathological changes such as inflammation, necrosis, apoptosis, or fibrosis observed in the liver tissue of SEAC treated groups. Moreover, the histological structure was very similar to that of the vehicle treated male and female ICR mice (Figure 2). Therefore, these results suggest that SEAC administration does not induce hepatotoxicity in the liver of male and female ICR mice.
Effects of SEAC on nephrotoxicity of male and female ICR mice

Finally, the nephrotoxicity against the treatment of SEAC was investigated in male and female ICR mice using serum biochemical and histological analyses of the kidney. BUN and Cr were maintained at constant levels between the vehicle treated group and the SEAC treated group without any significant alteration (Table 3). In addition, no specific pathological symptoms including degeneration and necrosis of the glomerulus and renal tubes were observed in the kidney tissue of any SEAC treated groups (Figure 3). Taken together, these results suggest that SEAC treatment for 14 days does not induce nephrotoxicity in male or female ICR mice.
Discussion

There are still many issues that must be overcome to successfully develop new drugs, including the large-scale procurement of sufficient source materials for bulk production, the potency and inherent toxicity of many natural products, and the development of suitable vehicles and dosing schedules for the administration of new drugs to clinical patients [11], although natural products have long been contributed as a source of leads.

### Table 1. Alteration of body and organ weights of female and male ICR mice treated with saponin-enriched extract of *A. cochinchinensis* (SEAC) for 14 days

| Category       | Vehicle | SEACL | SEACM | SEACH |
|----------------|---------|-------|-------|-------|
| **Male**       |         |       |       |       |
| Body weight (g) | 35.7±1.030 | 35.8±1.327 | 36.8±2.804 | 36.7±0.927 |
| Testis (g)     | 0.201±0.030 | 0.226±0.028 | 0.241±0.010 | 0.232±0.013 |
| Kidney (g)     | 0.545±0.064 | 0.557±0.074 | 0.581±0.026 | 0.56±0.018 |
| Spleen (g)     | 0.125±0.010 | 0.131±0.013 | 0.147±0.018 | 0.148±0.008 |
| Liver (g)      | 1.905±0.112 | 2.043±0.257 | 2.173±0.296 | 2.094±0.164 |
| Thymus (g)     | 0.038±0.007 | 0.043±0.006 | 0.049±0.008 | 0.046±0.006 |
| Heart (g)      | 0.19±0.016  | 0.188±0.020 | 0.209±0.032 | 0.203±0.012 |
| Lung (g)       | 0.217±0.010 | 0.215±0.013 | 0.230±0.014 | 0.236±0.003 |
| **Female**     |         |       |       |       |
| Body weight (g) | 29.8±0.678 | 30.9±1.428 | 30.6±1.855 | 29.2±1.122 |
| Ovary (g)      | 0.033±0.008 | 0.031±0.005 | 0.028±0.004 | 0.026±0.005 |
| Kidney (g)     | 0.311±0.009 | 0.333±0.011 | 0.338±0.019 | 0.337±0.019 |
| Spleen (g)     | 0.130±0.009 | 0.145±0.008 | 0.147±0.022 | 0.137±0.005 |
| Liver (g)      | 1.44±0.277  | 1.585±0.074 | 1.405±0.109 | 1.534±0.136 |
| Thymus (g)     | 0.084±0.013 | 0.092±0.022 | 0.062±0.009 | 0.076±0.012 |
| Heart (g)      | 0.132±0.007 | 0.134±0.006 | 0.139±0.013 | 0.134±0.018 |
| Lung (g)       | 0.27±0.012  | 0.283±0.024 | 0.262±0.014 | 0.241±0.011 |
| Brain (g)      | 0.506±0.031 | 0.51±0.014  | 0.519±0.010 | 0.515±0.020 |

The data shown represent the means±SD of three replicates.

### Table 2. Alteration on the urine parameters in male and female ICR mice after saponin-enriched extract of *A. cochinchinensis* (SEAC) treatment

| Category       | Vehicle | SEACL | SEACM | SEACH |
|----------------|---------|-------|-------|-------|
| **Male**       |         |       |       |       |
| Bilirubin (µmol/L) | 10.60±2.0 | 10.60±3.20 | 12.20±3.92 | 13.00±4.00 |
| Urobilinogen (µmol/L) | 36.00±24.49 | 31.98±11.45 | 33.00±8.00 | 36.00±24.49 |
| Leucocytes (WBC/µL) | 43.75±24.21 | 47.22±24.85 | 44.29±27.05 | 37.50±21.65 |
| Protein (g/L)   | 3.60±3.32  | 5.00±4.15  | 4.60±4.41  | 3.20±3.49  |
| pH              | 7.60±0.20  | 7.30±0.24  | 7.10±0.037 | 7.30±0.24  |
| Specific gravity| 1.02±0.00  | 1.03±0.00  | 1.02±0.01  | 1.03±0.00  |
| Ketones (mg/dL) | 1.00±0.00  | 1.00±0.00  | 1.00±0.00  | 1.00±0.00  |
| **Female**     |         |       |       |       |
| Bilirubin (µmol/L) | 11.67±3.77 | 11.67±3.77 | 9.00±0.00  | 10.00±3.77 |
| Urobilinogen (µmol/L) | 32.67±23.57 | 49.33±23.57 | 41.00±25.00 | 40.00±22.00 |
| Leucocytes (WBC/µL) | 58.33±23.57 | 62.50±21.65 | 44.29±27.05 | 54.29±22.05 |
| Protein (g/L)   | 1.00±0.00  | 0.83±0.30  | 1.67±0.94  | 3.20±0.30  |
| pH              | 8.00±0.00  | 8.00±0.00  | 7.38±0.48  | 7.30±0.24  |
| Specific gravity| 1.02±0.00  | 1.02±0.00  | 1.02±0.01  | 1.02±0.02  |
| Ketones (mg/dL) | 1.00±0.00  | 0.75±0.25  | 0.94±0.17  | 0.86±0.25  |

The data shown represent the means±SD of three replicates.
Toxicity of A. cochinchinensis extract

**Table 3.** Alteration on the serum parameters in male and female ICR mice after saponin-enriched extract of *A. cochinchinensis* (SEAC) treatment

|       | Category | Vehicle | SEACL | SEACM | SEACH |
|-------|----------|---------|-------|-------|-------|
| Male  | ALT (U/L)| 31.20±3.19 | 31.40±3.83 | 28.25±4.15 | 38.67±5.25 |
|       | AST (U/L)| 77.60±7.96 | 74.60±7.58 | 64.00±11.70 | 76.50±9.86 |
|       | ALP (U/L)| 74.80±12.73 | 87.00±17.76 | 90.67±15.41 | 78.75±16.98 |
|       | LDH (U/L)| 832.97±76.07 | 741.35±80.00 | 858.67±225.37 | 878.55±99.45 |
|       | BUN (mg/dL)| 26.40±4.57 | 26.37±5.11 | 29.41±3.06 | 28.34±4.73 |
|       | Cr (mg/dL)| 0.28±0.10 | 0.28±0.04 | 0.22±0.07 | 0.20±0.00 |
| Female| ALT (U/L)| 31.60±7.96 | 28.00±2.55 | 28.00±2.10 | 28.80±3.66 |
|       | AST (U/L)| 88.60±9.20 | 80.75±10.62 | 78.80±6.14 | 81.6±8.01 |
|       | ALP (U/L)| 117.25±23.97 | 100.00±2.83 | 100.60±17.43 | 108.40±22.81 |
|       | LDH (U/L)| 1.23±0.39 | 0.87±0.31 | 0.81±0.17 | 0.81±0.34 |
|       | BUN (mg/dL)| 31.08±4.23 | 27.33±2.70 | 28.40±1.88 | 28.46±2.80 |
|       | Cr (mg/dL)| 0.37±0.05 | 0.30±0.00 | 0.34±0.08 | 0.36±0.08 |

The data shown represent the means±SD of three replicates.

**Figure 2.** Effects of SEAC on liver toxicity in ICR mice. Liver tissues of male and female ICR mice were stained with H&E and histological structure was viewed at 400x magnification.
during the development of therapeutic drugs available for the treatment of human diseases [12,13]. It is very important to consider toxicological problems when developing novel drugs for treatment of human disease and to extend the therapeutic potential of conventional drugs [14]. In this study, we investigated the hepatotoxicity and nephrotoxicity of SEAC in ICR mice after short term administration to verify the possibility of using SEAC for clinical study because therapeutic effects of SEAC have been reported in inflammatory disease. The results presented here are the first to demonstrate that treatment with SEAC did not induce

**Figure 3.** Effects of SEAC on the kidney toxicity in ICR mice. Kidney tissues of male (A) and female (B) ICR mice were stained with H&E, and histological structure was viewed at 400× magnification.
significant toxicity against the livers and kidneys of male and female ICR mice at less than 600 mg/kg body weight.

Saponins are well known secondary metabolites of glycosidic nature that are abundantly distributed in higher plants, as well as some animal sources [15,16]. These compounds show varying ranges of pharmacological and biological activities such as anti-inflammatory, expectorant, vasoprotective, hypcholesterolemic, immunomodulatory, hypoglycemic, molluscicidal, anti-fungal, anti-parasitic and many others activities, while they also used for adjuvant production [17-19]. Despite their beneficial effects against clinical diseases, several studies have reported toxicity of saponin mixtures and saponin-enriched extracts. The oral administration (100 and 200 mg/kg) of water extract of saponins from *Argania spinose* for 3 months induced increased blood Cr levels and renal pathology without other pathological alterations [20]. However, treatment with a saponin mixture isolated from *Schefflera leucaentha* Viguier resulted in no significant differences in the histopathological structure of a control group and treated group, although BUN, Cr, AST, ALT and ALP levels were slightly increased in the group treated with the saponin mixture [21]. Furthermore, lambs treated with saponin-containing *Yucca schidigera* Roetzl juice (63 and 126 mg saponin/kg) showed acute tubular necrosis, dehydration and watery content in the gastrointestinal tract, as well as increased serum Cr and urea levels [22]. A 13 week subchronic toxicity study revealed that the lowest-observed-adverse-effect level (LOAEL) of saponin-rich and isoflavones-containing soybean extract was 707.2 and 751.8 mg/kg body weight/day, respectively [23]. In the present study, we investigated the heptatoxicity and nephrotoxicity of ICR mice treated with 150 (8.5 mg/kg of saponin), 300 (17 mg/kg of saponin) and 600 mg/kg (34 mg/kg of saponin) of SEAC. No significant toxicity was detected based on the body or organ weight, urine parameters, serum parameters or histological structure. These differences may have occurred because the concentration of saponins administered to animals was higher in most of the studies referenced above (>63 mg) than in the present study (34 mg).

The liver is well known as a primary target responsible for metabolism of various compounds including natural products, chemicals, drugs and alcohol, food additives and pesticides [24]. Some toxicity markers including the degree of steatosis, lobular inflammation, hepatocellular ballooning, and fibrosis can be detected during evaluation of liver toxicity caused by various compounds [25]. Among these, alterations in the levels of ALT, AST, ALP and LDH have been widely applied to measure liver toxicity [26-30]. Following treatment with saponin mixture isolated from *Schefflera leucaentha* Viguier, the level of AST, ALT and ALP increased [21]. However, the AST levels were slightly decreased in male rats treated with 2.5 and 5% of saponin-enriched and isoflavones-containing soybean extract, while ALT and ALP were maintained at constant levels [23]. Our results were very similar to those of a study in which animals were treated with soybean extract, although there are a few difference in the AST level.

In toxicological studies, the kidney is commonly considered an indicator organ for toxicity [31]. Generally, kidney dysfunction is caused by cellular damage induced by metabolites of the toxic chemicals that are excreted from the kidneys [32], and the severity of such damage is evaluated based on increases in red blood cells in the glomerulus and tubule luminal diameter upon histological analysis [33]. Other key markers for nephrotoxicity are serum levels of BUN and Cr that were dramatically increased in patients with renal trauma [34]. Therefore, the above parameters have been widely applied to evaluate kidney toxicity during preclinical and clinical experiments [35]. Moreover, in previous studies, treatment with high concentrations saponin and saponin mixtures induced significant alterations in the kidney histological structure and the level of serum markers [36]. Following saponin administration (200 mg/kg), increased Cr levels and deterioration of renal tubules were observed [20,37]. Moreover, the levels of BUN and Cr increased in groups treated with saponin mixture isolated from *S. leucaentha* Viguier (1,000 mg/kg for 14 days), saponin-containing *Y. schidigera* Roetzl juice (3,000 mg/kg for 21 days) and saponin rich soybean extract (5% of diet for 13 weeks) [21-23]. In the present study, no significant alterations in kidney weights or histopathological structure were detected throughout the experiment. Additionally, the levels of BUN and Cr remained constant in the serum of ICR mice treated with SEAC for 7 days. Our results differ from those of previous studies showing kidney toxicity of saponin related mixtures. These difference can be caused by differences in the dose of saponin containing mixtures applied to animals.

Taken together, the results of the present study suggest that SEAC with high antioxidant activity has no toxic effects on liver and kidney tissues of male and female ICR
mice. Furthermore, these findings provide vital information regarding the development of SEAC as a therapeutic drug for the treatment of various inflammatory diseases.

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Conflict of interests  The authors declare that there is no financial conflict of interests to publish these results.

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