Anti-oxidative and anti-inflammatory effects of spirulina on rat model of non-alcoholic steatohepatitis

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The pathogenesis of nonalcoholic steatohepatitis (NASH) remains unclear, but accumulating data suggest oxidative stress and the relationship between inflammation and immunity plays a crucial role. The aim of this study is to investigate the spirulina, which is a blue-green algae rich in proteins and other nutritional elements, and its component-phycocyanin effect on a rat model of NASH. NASH model rats were established by feeding male Wistar rats with choline-deficient high-fat diet (CDHF) and intermittent hypoxemia by sodium nitrite challenge after 5 weeks of CDHF. After experimental period of 10 weeks, blood and liver were collected to determine oxidative stress injuries and efficacies of spirulina or phycocyanin on NASH model rats. In the NASH model rats, increase in plasma liver enzymes and liver fibrosis, increases in productions of reactive oxygen species from liver mitochondria and from leukocytes, the activation of nuclear factor-kappa B, and the change in the lymphocyte surface antigen ratio (CD4+/CD8-) were observed. The spirulina and phycocyanin administration significantly abated these changes. The spirulina or phycocyanin administration to model rats of NASH might lessen the inflammatory response through anti-oxidative and anti-inflammatory mechanisms, breaking the crosstalk between oxidative stress and inflammation, and effectively inhibit NASH progression.

Key Words: non-alcoholic steatohepatitis, anti-oxidant, oxidative stress, anti-inflammation, spirulina

Non-alcoholic fatty liver disease has become prevalent in both developed and developing countries. Although the condition is usually considered benign, it can progress to non-alcoholic steatohepatitis (NASH), which may then lead to cirrhosis, hepatocellular carcinoma and liver failure.(1)

NASH is a progressive liver disorder that occurs in patients without significant alcohol consumption, and, histologically, it resembles the alcoholic liver disease with macrovesicular steatosis, spotty necrosis, inflammation and fibrosis.(2,3) The “two-hit theory” provides the most widely accepted explanation describing the progression of NASH. This hypothesis states that hepatic steatosis marks the first stage of NASH development. The “first hit” is considered to be related to the inflow and the increased synthesis of fatty acids, dysfunction of β-oxidation and obstruction of the excretion of very low density lipoprotein causing the excessive accumulation of fatty acids and neutral fat in liver hepatocytes. The “second hit” has been considered to be related to the oxidative stress caused by excess generations of reactive oxygen species (ROS) and free radicals, elevation of pro-inflammatory cytokines and appearance of mitochondrial dysfunction, all of which lead to liver inflammation and fibrosis.(4–6)

Here we describe a new and more clinically-relevant animal model of NASH (Patent application No. PCT/JP2007/52477), and demonstrated that oxidative stress causes extensive hepatic fibrosis and cirrhosis in this model rats.(7) To our knowledge, there are few published studies that have investigated the potential efficacy of antioxidant supplements against NASH progression. Information described in reports published by other groups, together with those presented in our previous papers, encourage us to select an excellent antioxidant that could potentially restrain the progression of NASH.(8–10)

Spirulina (SP, Spirulina platensis) is blue-green algae, contains 62% amino acids and is the world’s richest natural source of vitamin B12 and contains a whole spectrum of natural mixed carotenoids and xanthophyll phytopigments. SP is wrapped with a soft cell wall formed from complex sugars and proteins. Actually it is gaining more attention because of its nutritional and various medicinal properties.(11,12) SP is used as a food supplement and the nutritional and therapeutic values have been well documented.(13) SP has been claimed to possess immunomodulatory,(14) antioxidant,(15,16) anticancer,(17,18) antihyperlipidemic(19,20) and anti-diabetic properties(21) and shown to prevent lead toxicity.(22,23)

Phycocyanin (PC), one of the major biliproteins present in SP, is often used as a dietary nutritional supplement in many countries due to its therapeutic value, including hepatoprotective, anti-coagulant, neuroprotective and ROS-scavenging actions.(13,23–27) In addition, the anti-inflammatory activity of PC has been demonstrated in various in vitro studies and in vivo experimental models such as mice with arthritis or sepsis.(28–30)

As the antioxidative and anti-inflammatory properties of SP were expected to be able to halt the NASH progression, the present study was aimed to investigate the efficacies of SP and PC, and to elucidate their mechanisms.

Materials and Methods

Animals. Male Wistar rats (Shimizu Experimental Animals, Shizuoka, Japan), weighing 160–170 g and six weeks of age were used in this study. They were housed in the Animal Research Center of Okayama University in a temperature-controlled room (22 ± 1°C) with a relative humidity of 50 ± 10% and a 12 h light/dark cycle (lights from 08:00 to 20:00). This study was performed in accordance with the Ethics Review Committee for Animal Research of the Okayama University. 

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**Experimental design (Fig. 1).** The rats were fed either standard chow (control group, \( n = 8 \)) for rodents (MF laboratory chow, Oriental Yeast Co., Tokyo, Japan) or choline-deficient high-fat (CDHF) diets (Oriental Yeast Co., \( n = 48 \)) through the experiment period of 10 weeks. Fatty liver was induced by the feeding CDHF for 4 weeks. In the fifth week, rats on CDHF were divided randomly into six groups to equalize their body weight. In the following experiment period of 10 weeks, fatty liver was induced by the feeding CDHF for 4 weeks. In the fifth week, rats on CDHF were divided randomly into six groups to equalize their body weight. The remaining rats were divided randomly into six groups to equalize their body weight.

- The CDHF group received CDHF diet alone for 16 weeks. The NASH groups were added with injections of sodium nitrite, 50 mg/kg/day i.p., for the subsequent 6 weeks at the 10th week of continuous CDHF diets. In the NASH + 2SP and NASH + 6SP groups, spirulina, 2 g or 6 g/kg/day, was given p.o. concurrently during the period of CDHF diet and nitrite injection. In the NASH + 0.4PC and NASH + 1.2PC, the PC administration of 0.4 g or 1.2 g/kg/day was performed concurrently during the period of CDHF diet and nitrite injection.

**Blood and tissue preparation.** All work was performed in an aseptic environment. The rats were killed by anesthetizing with diethyl ether. Blood samples were collected by vena cava inferior puncture with syringe containing heparin, and whole body perfusion was performed to left ventricular with 0.1 M potassium phosphate buffer (pH 7.4) containing 0.1 M potassium chloride was mixed with 0.4% dodecyl maltoside, 5 mM benzamidine (Nacalai Tesque Inc., Kyoto, Japan) daily to induce methemoglobinemia (intermittent hypoxia stress) starting from the 5th week of CDHF for 6 weeks. In the NASH + 2SP group \( (n = 8) \) or the NASH + 6SP group \( (n = 8) \), SP at the dose of 2 g/kg/day or 6 g/kg/day, p.o. was given concurrently during the period of nitrite injection. In the NASH + 0.4PC group \( (n = 8) \) or the NASH + 1.2PC group \( (n = 8) \), PC at the dose of 0.4 g/kg/day or 1.2 g/kg/day, p.o. was given concurrently during the period of nitrite injection. The PC dosage was set according to the PC content of SP.

**Histopathological examination.** Paraffin-embedded tissues were sectioned at 4 \( \mu \)m thickness and stained with Masson trichrome for fibrosis. Histological assessment of tissue morphology was performed using an Olympus light microscope (Olympus, Tokyo, Japan).

**Plasma assay for liver enzymes.** Plasma alanine aminotransferase (ALT) and aspartate aminotransferase (AST) were determined with commercial kits (Wako Pure Chem. Ind. Ltd.).

**Mitochondrial protein concentration.** Protein concentration of the liver mitochondria was determined by Lowry method.

**Derivation of ROS from liver mitochondria.** The ROS derivation from liver mitochondria was estimated by electron spin resonance (ESR) spectrometer (JES-RE1X JEOL RE-1X, JEOL, Tokyo, Japan) connected with the data analyzer (WIN-RAD, Radical Research Inc., Tokyo, Japan), using 5,5-dimethyl-1-pyrroline-1-oxide (DMPO) (Labotech Co., Tokyo, Japan) as a radical-trapping reagent.

35 \( \mu \)L of mitochondria suspension in 0.25 M sucrose Tris-HCl buffer containing 0.1 M potassium chloride was mixed with 25 \( \mu \)L of substrate solution containing 0.4% dodecyl maltoside, 40 mM sodium glutamate, 40 mM disodium malate and 800 mM disodium succinate in 50 mM potassium phosphate buffer, pH 7.4, 20 \( \mu \)L of 4.6 M DMPO and finally 1.0 mM NADH (Oriental Yeast Co., Osaka, Japan), then was incubated at 37°C. After 5 min...
incubation, 45 μL aliquots of above-mentioned mixture were placed into a capillary glass tube, and this was in turn introduced into a quartz tube, and then measured at room temperature. The instrument conditions were the following: about 9.4 GHz with 100-kHz modulation frequency, central field ± sweep width of 336.0 ± 5 mT, microwave power of 8 mW, field modulation width of 0.1 mT, time constant of 0.1 s, sweep speed of 120 G/min. The magnetic field was calculated from the splitting of divalent manganese (Mn²⁺), in which the distance from the third to fourth signal is 8.69 mT.

In this measurement, the characteristic ESR spectrum of the spin adducts of DMPO and hydroxy radical (DMPO-OH) was detected. The ROS derivation was estimated by the ratio of the 2nd signal intensity of the DMPO-OH spectrum/the 3rd signal of Mn²⁺.

**Liver lipid peroxidation assay.** The liver lipid peroxidation was determined with chemiluminescence (CL) method. 140 μL of stored for lipid peroxidation supernatant samples were mixed with 20 μL of 130 μg/mL luminol (Wako Pure Chem. Ind. Ltd.) and 20 μL of Hanks’ Balanced Salt Solution (HBSS, pH 7.4), then was incubated and analyzed in the detector (TriStar LB941, Berthold Japan Co. Ltd., Tokyo, Japan) at 37°C for the baseline CL intensity. After 5 min incubation, 80 μL of 2.2, 5, tert-butyl hydroperoxide (tert-BuOOH; SIGMA, St. Louis, Mo) was added. CL intensity was detected from the cleavage of lipid peroxides induced by tert-BuOOH addition, for 120 min with incubating at 37°C. In this measurement, lipid peroxidation levels were estimated by the accumulated CL intensity by addition of tert-BuOOH subtract from the baseline CL intensity for 120 min.

**Leukocyte oxygen radical production.** 125 μL of 50-fold diluted whole blood samples were mixed with 25 μL of 300 μg/mL luminol and 20 μL of HBSS then was incubated at 37°C. After 5 min incubation, 80 μL of 0.0781 μg/mL phorbol 12-myristate 13-acetate (PMA; SIGMA) was be added. CL intensity was detected from the cleavage of lipid peroxides estimated by measuring the amount of oxidized luminol by oxygen free radicals after PMA stimulation, as the same manner mentioned above.

**Nuclear extract and Western blot analyses to determine nuclear transcription factors.** The nuclear fractions sample suspended in 50 mM HEPES buffer (pH 7.4) containing 0.1 M potassium chloride, 3 mM magnesium chloride, 1 mM ethylenediaminetetraacetic acid, 10% Glycerol, 0.1 mM phenylmethylsulfonyl fluoride, 5 μg/mL pepstatin A, 5 μg/mL leupeptin and 2 μg/mL aprotinin, and centrifuged at 22,000 × g for 20 min at 4°C. The supernatant was used as nucleoprotein samples. For protein quantification, Lowry method was used.

The nucleoprotein sample was diluted to 6 mg/mL, then mixed with sample buffer (62.5 mM Tris-HCl pH 6.8, containing 25% glycerol, 2% sodium dodecyl sulfate, 5% 2-mercaptoethanol, 0.01% bromophenol blue) and denatured at 95°C for 5 min. The samples (in 30 μg protein/10 μL) were separated on SDS-12.5% polyacrylamide gel (Bio-Rad Laboratories Inc., Berkeley, CA), then transferred to polyvinylidene fluoride (PVDF) membrane using a transblot apparatus (Bio-Rad Laboratories Inc.). The membranes were blocked in 5% nonfat milk dissolved in TBS-T buffer (25 mM Tris-HCl buffer, pH 7.4, containing 0.15 M sodium chloride and 0.1% Tween20) for 1 h at room temperature. The membranes were incubated with primary antibodies as follows: mouse monoclonal anti-rat nuclear factor-kappa B (NF-kB) (1:1000; Santa Cruz Biotechnology, Santa Cruz, CA) or rabbit polyclonal anti-rat Histone H1 (1:200; Santa Cruz Biotechnology) for 1 h. And then incubated with secondary antibodies for goat anti-mouse IgG-HRP (1:5000; Santa Cruz Biotechnology) or goat anti-rabbit IgG-HRP (1:5000; Santa Cruz Biotechnology) for 30 min. Protein bands were visualized using enhanced Chemiluminescence Luminol Reagent (Santa Cruz Biotechnology). The results were standardized by Histone H1.

**Preparation of the liver microsomal fraction.** Aliquots of liver microsomes (5 μg) were placed in sample buffer (62.5 mM Tris-HCl pH 6.8, containing 25% glycerol, 2% sodium dodecyl sulfate, 5% 2-mercaptoethanol, 0.01% bromophenol blue) and denatured at 95°C for 5 min. Samples were separated on SDS-12.5% polyacrylamide gel and transferred to PVDF membranes, blocked by Western blot analyses with the primary rabbit anti-human/rat cytochrome P450 enzyme (CYP2E1) polyclonal antibodies (1:1500; Chemicon International, Temecula, CA) or mouse monoclonal anti-rabbit β-actin (1:2000, Santa Cruz Biotechnology) for 1 h. Proteins were detected with peroxidase-conjugated secondary anti-rabbit IgG antibody or goat anti-mouse IgG HRP (1:5000; Santa Cruz Biotechnology) for 1 h and used enhanced Chemiluminescence Luminol Reagent. The results were standardized by Histone H1.

**Plasma myeloperoxidase (MPO) activity.** 225 μL of the plasma was mix with 315 μL of 10 mM citrate buffer, pH 5.0, then aliquots of 90 μL of these mixed liquid were pipetted into four wells. The 540 nm absorbance was monitored about the two wells to which 120 μL of cold stop solution (4 N sulfuric acid) was added to stop the reaction to detect the background absorbance. 30 μL of MPO substrate solution containing 6 mM 3, 5, 5′, 6-tetramethylbenzidine dihydrochloride, 120 μM resorcinol, and 4.4 mM hydrogen peroxide in distilled water was added to each well, and the reaction was stopped after 2 min with 120 μL of cold stop solution, subsequently measured by absorbance at 450 nm. As an additional control, 90 μL of dilution buffer (without liver extract) were pipetted into four wells, 30 μL of substrate buffer was added, followed by 120 μL of of stop solution after 2 min. No color reaction was observed in these control wells. The MPO activity of the liver sample was calculated by subtracting the mean background absorbance and is expressed as change in optical density per minute.

**Isolation of leukocytes and flow cytometry analysis of lymphocyte surface antigen CD4, CD8.** The peripheral blood was divided and erythrocytes were lysed with ammonium chloride solution, and washed twice in phosphate-buffered saline containing 0.1% sodium azide and 0.1% bovine serum albumin (BSA). The final pellet was resuspended in same buffer.

Liver tissue was dissected into 1 mm³ fragments in HBSS medium containing 2% fetal bovine serum (FBS), 25 mM HEPES, 0.6% BSA, 0.125% type IV collagenase and 0.002% deoxyribonuclease I (DNase I) to each sample and the mixtures incubated with gentle agititation at 37°C for 30 min. Undissociated tissue was removed by filtration through a 30 μm nylon mesh and the cells were washed twice with HBSS containing 0.001% DNase I and 25 mM HEPES with centrifugation at 250 × g for 10 min. The hepatocyte-rich matrix was removed by centrifugation for 1 min at 30 × g and the final pellet was resuspended in phosphate-buffered saline containing 0.1% sodium azide and 0.1% BSA.

For flow cytometry, the cells were incubated with fluorescein isothiocyanate (FITC) or phycoerythrin (PE)-conjugated monoclonal antibodies (antimouse CD4 [FITC], or antimouse CD8 [PE]) (all sourced from eBioscience, San Diego, CA) for 20 min, washed twice times. Then cell fixation using 400 μL of 1% paraformaldehyde immediately before determination. At least 10,000 cells in fluorescent channel 1 for CD4, and fluorescent channel 2 for CD8 were detected using FACSCalibur™ flow cytometer (BD Biosciences, Franklin Lakes, NJ). Results were expressed as mean fluorescence intensity for a given molecule per cell.

**Statistical analyses.** Quantitative data were expressed as the mean ± SEM of six rats. Differences among groups were examined by one-way analysis of variance (ANOVA) followed by Tukey-Kramer multiple comparison test. A value of p<0.05 was considered statistically significant.
Results

Histological evaluation of experimental liver samples.
The gross changes of experimental liver samples were shown in Fig. 2. The CDHF rat livers (Fig. 2b) were significantly enlarged compared to Control rats (Fig. 2a). Severe liver atrophy and cirrhosis were observed in NASH livers (Fig. 2c). The administration of SP or PC improved these changes (Fig. 2 d–g).

Masson trichrome stain was used to see fibril formation. The macrovesicular steatosis was remarkable in all zones of the liver lobe in CDHF rats (Fig. 3b). In the NASH rats, steatosis, liver fibrosis, and necrosis were quite advanced. The pseudolobule was also observed (Fig. 3c). The administration of SP or PC abated the state of the NASH (Fig. 3 d–g).

Effects of SP or PC on plasma hepatic enzyme activities.
The plasma levels of AST and ALT were significantly higher in

Table 1. Effects of SP or PC on body weight and liver index

<table>
<thead>
<tr>
<th>Group</th>
<th>Body weight (g)</th>
<th>Liver wet weight (g)</th>
<th>Liver index (%)</th>
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<tbody>
<tr>
<td>Control</td>
<td>366.77 ± 26.78</td>
<td>10.56 ± 1.93</td>
<td>2.88</td>
</tr>
<tr>
<td>CDHF</td>
<td>225.33 ± 25.74**</td>
<td>11.49 ± 1.90</td>
<td>5.10**</td>
</tr>
<tr>
<td>NASH</td>
<td>218.33 ± 24.35**</td>
<td>7.35 ± 2.16</td>
<td>3.33</td>
</tr>
<tr>
<td>NASH + 2SP</td>
<td>248.13 ± 30.94**</td>
<td>12.96 ± 3.41</td>
<td>5.22</td>
</tr>
<tr>
<td>NASH + 6SP</td>
<td>312.75 ± 20.25**</td>
<td>9.54 ± 2.02</td>
<td>3.05**</td>
</tr>
<tr>
<td>NASH + 0.4PC</td>
<td>222.33 ± 12.23**</td>
<td>8.58 ± 1.84</td>
<td>3.84</td>
</tr>
<tr>
<td>NASH + 1.2PC</td>
<td>221.67 ± 22.25**</td>
<td>8.57 ± 2.10</td>
<td>3.82</td>
</tr>
</tbody>
</table>

Each value denotes the mean ± SEM of 8 rats. **p<0.01 vs Control group; **p<0.01 vs NASH group. Liver index (%): liver wet weight/body weight ×100.

Fig. 2. Macroscopic views of experimental rat livers at the end of the study. See the Fig. 1 for experimental details. (a) The Control group, (b) CDHF group, (c) NASH groups, (d) NASH + 2SP group, (e) NASH + 6SP group, (f) NASH + 0.4PC group and (g) NASH + 1.2PC group, Scale bar = 500 μm.

Fig. 3. Microscopic pictures of experimental rat livers. The experimental design is the same with Photo 1. Groups: (a) Control, (b) CDHF, (c) NASH, (d) NASH + 2SP, (e) NASH + 6SP, (f) NASH + 0.4PC and (g) NASH + 1.2PC, Scale bar = 500 μm.
NASH rats compared with rats of the Control group ($p<0.01$, Fig. 4). Except NASH + 0.4PC group, the administration of SP or PC to NASH rats significantly improved the AST or ALT levels compared with NASH group ($p<0.05$, $p<0.01$).

**Effects of SP or PC on mitochondrial ROS derivation.** ROS leakage from mitochondria was evaluated by spin trapping method of ESR spectroscopy, the manganese signal as an internal standard. The amount of ROS derived from mitochondria was expressed numerically by measuring the relative peak heights of the second peaks from the DMPO-OH signals (Fig. 5). We found the highest signal intensities were obtained in NASH group. SP or PC administration caused a significant decrease in detectable levels of liver mitochondrial ROS generation compared with the NASH group ($p<0.05$, $p<0.01$).

**Effects of SP or PC on liver lipid peroxidation.** The liver lipid peroxidation levels were significantly higher in NASH group, compared with the control group ($p<0.01$, Fig. 6). Administration of SP or PC was significantly prevented the lipid peroxidation of NASH rats ($p<0.05$).

**Effects of VCPL on CYP2E1 induction in the liver microsomes.** The CYP2E1 expression in the NASH group was significantly higher than the control group ($p<0.01$, Fig. 7). However, SP or PC administration did not change the CYP2E1 expression for NASH.

**Effects of SP or PC on leukocyte ROS production.** In order to assess the state of leukocyte priming, PMA-stimulated ROS productions from leukocytes were measured (Fig. 8). In this study, CDHF and NASH groups showed significantly higher oxygen radicals signal from leukocyte compared with rats of the Control group ($p<0.01$). SP or PC administration, significantly suppressed ROS produced from leukocytes ($p<0.05$ or $p<0.01$).

**Effects of SP or PC on NF-$\kappa$B activation.** NF-$\kappa$B, a transcription factor linking between oxidative stress and inflammation, migrates to the nucleus and regulate inflammatory protein transcription when cells are activated by oxidative stress. The increase in the NF-$\kappa$B import into nuclei of NASH rats ($p<0.01$, Fig. 9) is significant. SP or PC administration, significantly reduced the activation of NF-$\kappa$B ($p<0.05$ or $p<0.01$).

**Effects of SP or PC on neutrophil activation.** Fig. 10 showed that the plasma MPO activity was significantly higher in NASH group than compared with the control group ($p<0.01$). Administration of SP significantly decreased plasma MPO activity ($p<0.05$ or $p<0.01$).
Effects of SP or PC on Lymphocyte CD4⁺/CD8⁺ ratios.
The lymphocytes CD4⁺/CD8⁺ ratio of peripheral blood was significantly higher in NASH group, compared with the control group (p<0.05, Fig. 11A), suggesting the hyperactivity of immune system in NASH rats. SP or PC administration significantly decreased the ratio of CD4⁺/CD8⁺ on peripheral blood (p<0.05 or p<0.01). Lymphocytes CD4⁺/CD8⁺ ratio in liver showed no significant difference, but showed similar tendency in the peripheral blood (Fig. 11B).

Discussion

Recently, NASH is a very popular chronic liver disease worldwide, and it is the major cause of the elevated liver enzymes (ALT and AST in plasma) in developed countries. A major problem of NASH is the lack of suitable experimental animal models. Our NASH model was developed on the basis of “Two hit theory”, and can reproduce the typical hepatic lesions of NASH, namely, steatosis, inflammation, and early fibrosis. Our new rat model of NASH might be useful for understanding the disease treatment and prevention and we suggest its application as an excellent model for further mechanistic insight into NASH pathogenesis.

In the present study, we showed that administration of SP or PC improved liver atrophy, liver fibrosis and liver function (Fig. 2 and 4). We also showed that oral administration of SP or PC decreased the activities of AST and ALT in NASH rats. In a word, SP or PC successfully slowed down the development of steatohepatitis via histopathological and biochemical improvements. These effects suggest that SP or PC could play the role in protecting body against NASH.

Although the reasons for the deleterious effects of steatosis are still incompletely understood, there is growing evidence that, in the presence of oxidizable fat in the liver, ROS induces depletion of mitochondrial DNA, up-regulation of uncoupling protein 2, reduced activity of respiratory chain complexes, and impaired mitochondrial β-oxidation, resulting in mitochondrial dysfunction and abnormal mitochondrial morphology. ROS triggered lipid peroxidation generates several reactive aldehydes. ROS and reactive aldehydes, in turn, further impair mitochondrial function, to further increase mitochondrial ROS derivation. A ROS-dependent...
vicious circle may thus ensue. Therefore, the mitochondria are the most important intracellular source of ROS in NASH.\(^{(44–47)}\) Our data also showed that the increase ROS derivation from mitochondrial and higher lipid peroxidation levels in NASH model rats liver (Fig. 5 and 6).

Some of the active constituents of SP have been reported to possess strong activity for provoking free radical scavenging enzyme system.\(^{(16,38–40)}\) SP or PC administration in our NASH model system improved pathological changes, and this suggests that SP or PC supplementation could prevent excessive production of ROS, to regulate the balance of pro- and anti-oxidants. We, then, tried to elucidate the mechanism of SP or PC action.

CYP2E1 is a well-known other sources of oxidative stress and its levels are elevated in the liver of NASH patients.\(^{(41)}\) Recent studies indicated that CYP2E1 plays a key role in the pathogenesis of NASH with its ability to initiate oxidative stress and increased lipid peroxidation.\(^{(42–45)}\) Western blot analysis was used to examine the induction of CYP2E1 expression with the development of our NASH model. Our data showed higher CYP2E1 expression in the NASH model group, but SP or PC administration could not resolve these changes (Fig. 7), and higher CYP2E1 expression in our model is not associated with the progression of the disease.

It was also proposed that pro-inflammatory condition was the key predictor of eventual histological progression in NASH and could be a potential major therapeutic target.\(^{(44–46)}\) Inflammation causes neutrophil activation and infiltration into the damaged tissues. The neutrophil infiltration may also be an important source of various pro-inflammatory mediator productions, including cytokines and oxygen-derived free radicals.\(^{(47)}\) We observed that PMA stimulated ROS production from leukocytes were measured (Fig. 8) and SP or PC administration inhibited this inflammatory status.

NF-κB is known to be ubiquitously expressed and to play a major role in controlling the expression of protein involved in immune, inflammatory and acute phase response. Without stimulation, NF-κB is in an inactive state bound to its inhibitor IκB in the cytoplasm. Various agonists, such as IL-1, TNF-α and TLR ligands, activate NF-κB. Then the NF-κB undergoes nuclear translocation, where it binds to and stimulates transcription of target genes.\(^{(48)}\) In this paper, the western blot analysis showed that the activation of the NF-κB was involved in the development of NASH (Fig. 9). The increased in MPO activity, an indicator of neutrophil activation and an important enzyme involved in the generation of reactive oxygen species,\(^{(49)}\) was observed in NASH model (Fig. 10). SP or PC administration significantly reduced these changes, suggesting that SP could suppress NF-κB nuclear transport to prevent pro-inflammatory transcription and inhibit neutrophil activation.

Many of the current data show that innate immune processes both within and outside the liver are involved in NASH.\(^{(50,51)}\) The percentages of CD4⁺ lymphocytes and CD8⁺ T lymphocytes and the ratio of CD4⁺/CD8⁺ conveniently estimate the immune state. In this study, we measured the differentiation antigens of CD4⁺ and CD8⁺ in peripheral blood of NASH rats by flow cytometry to further determine the effects of SP on cellular immune function (Fig. 11). The data show that the SP can effectively normalize the immune imbalance.

In our NASH model, NaNO₂ was injected intraperitoneally, to provoke methemoglobinemia and decay the oxygen transport ability, which induced the intermittent hypoxemia. The enhanced oxidative stress was also demonstrated in NASH rats. So, it is firmly expected the alteration of heme oxygenase in our NASH rats. Incidentally there is indicated that Bach1 ablation exerts hepatoprotective effect against steatohepatitis presumably via heme oxygenase induction and may be a potential therapeutic target.\(^{(52)}\) SP efficacy would associate with the expression of Bach1 or heme oxygenase.

The present study exhibited that SP could prevent the excessive ROS derivation from the liver mitochondrial energy metabolism. Furthermore, we demonstrated that SP could relieve the ROS production from irritable inflammatory cells and the alteration of surface antigen expression of lymphocytes in the peripheral blood of NASH rats, as protective factors in liver inflammation and fibrosis. Thus, we are convinced SP also may work against NASH through anti-inflammatory and immuno-modulatory effects.

Lastly, it is well known that disturbance of lipid metabolism is a basic pathogenesis of NASH. The present study exhibited the SP efficacy against hepatic fat accumulation. Therefore, besides the anti-oxidant and anti-inflammatory activities of SP, possible metabolic roles or other mechanisms of SP in NASH rats are also considered, such as the enterobacterial flora improvement. It may play a role in ameliorating fatty liver, as the association with gut probiotic flora and attenuation of hepatic fat accumulation has been demonstrated,\(^{(53)}\) besides reducing an inflammatory signal. However, the effect of SP on adipose metabolism needs further research.

In conclusion, we demonstrated in the present study that the anti-oxidant and anti-inflammatory activities of SP or PC can prevent the progression of NASH by normalizing the redox balance disorder and inflammatory reaction. These results not only deepen our understanding of SP, but also strongly imply that SP can be potentially used as a promising agent for prevention and therapy of NASH in clinical trials.

Acknowledgments

We would like to express the appreciation to Pharm. M. Azusa Hasegawa and Pharm. M. Shiho Sugimoto for their kind assistance.

Abbreviations

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<th>Abbreviation</th>
<th>Description</th>
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<tr>
<td>ALT</td>
<td>Alanine aminotransferase</td>
</tr>
<tr>
<td>AST</td>
<td>Aspartate aminotransferase</td>
</tr>
<tr>
<td>BSA</td>
<td>Bovine serum albumin</td>
</tr>
<tr>
<td>CDHF</td>
<td>Cholinedeficient high fat</td>
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<tr>
<td>CL</td>
<td>Chemoluminescence</td>
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<tr>
<td>DMPO</td>
<td>5,5-Dimethyl-1-pyrrrole N-oxide</td>
</tr>
<tr>
<td>FITC</td>
<td>Fluorescein isothiocyanate</td>
</tr>
<tr>
<td>MPO</td>
<td>Myeloperoxidase</td>
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<tr>
<td>NASH</td>
<td>Nonalcoholic steatohepatitis</td>
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<tr>
<td>NF-kB</td>
<td>Nuclear factor-kappa B</td>
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<tr>
<td>PC</td>
<td>Phycocyanin</td>
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<tr>
<td>PE</td>
<td>Phycocerythrin</td>
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<tr>
<td>PMA</td>
<td>Phorbol 12-myristate 13-acetate</td>
</tr>
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<td>PVDF</td>
<td>Polyvinylidene fluoride</td>
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<tr>
<td>ROS</td>
<td>Reactive oxygen species</td>
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<td>Spirulina</td>
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<td>tert-BuOOH</td>
<td>tert-butyl hydroperoxide</td>
</tr>
</tbody>
</table>

Conflict of Interest

The authors declare no conflicts of interest that are relevant to this article.


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