Chemopreventive effects of fermented brown rice and rice bran against 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone-induced lung tumorigenesis in female A/J mice

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Abstract. The most common cause of human lung cancer is suggested to be exposure to the carcinogens in tobacco smoke. Among the multiple chemicals in tobacco smoke, 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone (NNK) has been regarded as one of the important agents for generation of lung cancers. Previously, our studies proved that fermented brown rice and rice bran (FBRA) has chemopreventive effects against carcinogenesis of the colon, liver, stomach, bladder and esophagus. In the present study, we examined possible chemopreventive effects of FBRA on the NNK-induced lung tumorigenesis in mice. Six-week-old female A/J mice were divided into 8 groups, and groups 1-5 were given NNK (10 mmol) by i.p. injection at week 7. Groups 2 and 3 were fed with diet containing 5 and 10% FBRA during the initiation phase, respectively. Groups 4 and 5 were fed with 5% and 10% FBRA during the post-initiation phase. Groups 1 and 6 were given control diet throughout the experiment. Groups 7 and 8 were given the diet containing 5 and 10% FBRA throughout the experiment, respectively. In both initiation (group 3) and post-initiation phase (group 5), 10% FBRA exposure significantly reduced the multiplicity of lung tumor (group 3, 2.35±2.13; group 5, 3.00±1.52; group 1, 4.08±1.85; p<0.006 and 0.04, respectively). Furthermore, administration of FBRA during the post-initiation phase significantly decreased the tumor size in comparison with that of control mice (0.66±0.32 vs. 0.77±0.33 mm). Treatment of 10% FBRA significantly reduced the mRNA expression levels of cytochrome P450 2A5 (Cyp2a5), which is known to be closely related to the human CYP2A6 enzyme that is involved in the mutagenic activation of NNK, in the lung but not liver tissues. A significantly reduced index of Ki67 positivity of lung tumors in group 5 was confirmed when compared with tumors of the control group (0.065±0.016 vs. 0.114±0.025). These findings suggest that FBRA has inhibitory effects on NNK-induced pulmonary tumorigenesis in A/J mice in both during initiation and post-initiation treatment, which is possibly associated with the induction of Cyp2a5 in the lung and the reduced proliferation rate of tumor cells. FBRA may be a promising chemopreventive agent for human lung cancers.

Introduction

Cancer is a leading cause of death worldwide. Lung cancer is the most common cause of cancer death, with 1.18 million deaths (17.6%) of the world total exceeds stomach cancer (700,000 deaths) and liver cancer (598,000 deaths). However, there is limited information regarding effective agents for prevention of lung cancers. For the year 2000, about 85% of lung cancer in men and 47% of lung cancer in women is estimated to concern with tobacco smoking (1). According to the report by the International Agency for Research on Cancer (IARC), there are 55 carcinogens in tobacco smoke. Among the multiple ingredients in tobacco smoke, 20 carcinogens are likely to be involved in lung cancer induction in laboratory animals or humans. Of these, polycyclic aromatic hydrocarbons (PAHs) and a tobacco-specific nitrosamine 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone (NNK) are suggested to relate to carcinogenesis (2,3). NNK has been suggested to link to the etiology of lung cancer in humans as well as to the induction of tumors in commonly used rodents like rat, mouse and hamster, independently of its route of administration (4,5). NNK treatment in...
A/J mice results in the development of lung tumors that are similar to human adenocarcinoma, a representative type of human lung cancer in morphology and molecular characteristics (6-8).

Cytochrome P450 (CYP) is widely known as a family of enzymes metabolizing a variety of xenobiotics, including drugs and carcinogens. CYP plays central roles in the activation of many environmental chemicals to generate reactive intermediates (9). In vitro, enzymes from the CYP2A subfamily efficiently catalyze the metabolic activation of a number of nitrosamines, including NNK. However, the ability of each CYP2A enzyme to metabolize nitrosamines to their active forms is different from others (10,11). Human CYP2A6 enzyme is known to be responsible for the metabolic activation of chemicals including nicotine (12). Indeed, Japanese male smokers with a CYP2A6 gene deletion-type polymorphism have been shown to exhibit reduced lung cancer risk in a hospital-based case control study (13,14). In mice, function of Cyp2a5 enzyme is closely related to that of human CYP2A6 that is expressed in many tissues, including lung and liver (15).

One approach to restrain cancer incidence is a chemopreventive intervention through which the disease can be totally prevented, delayed or fixed dormant by one or more natural occurring and synthetic chemical agents. Rice is one of the major cereal foods eaten as a staple food worldwide, especially in Asian countries. Rice seeds and rice germ contain fiber and several kinds of antioxidants, such as ferulic acid (16), phytic acid (17), tocochromanols and oryzanol (18). Among them, fiber and ferulic acid have been reported to prevent chemically-induced aerodigestive tract carcinogenesis in animal models (19-21). Rice germ itself or the compounds contained in the rice bran or germ have chemopreventive effects against carcinogenesis in the colon (22), liver (23), stomach (24), esophagus (25) and bladder (26) of rodents. FBRA is a processed food prepared by fermented brown rice and rice bran with Aspergillus Oryzae. It is already known that FBRA acts as a potent free radical scavenger (27). The significance of brown rice and rice germ as modulators of carcinogenesis prompted us to explore FBRA for its potential anticancer properties. In this study, we examined the chemopreventive activity of FBRA on the tobacco-derived nitrosamine (NNK)-induced lung tumorigenesis in female A/J mice in initiation and post-initiation phases.

Materials and methods

Animals. Female A/J mice, 3 weeks of age, were obtained from Japan SLC, Inc. (Hamamatsu, Japan) and housed in wire cages (3 or 4/cage) under controlled conditions of humidity (50±10%), lighting (12-h light/dark cycle) and temperature (23±2°C), with free access to water and were bred and maintained on a basal diet, CE-2 (CLEA Japan Inc., Tokyo, Japan), until 6 weeks of age.

Chemicals. Powdered CE-2 diet was used as the basal diet throughout the study. The experimental diets were prepared by mixing 5.0 and 10.0% FBRA with CE-2 diet. FBRA was supplied by Gennai Koso Co., Ltd. (Sapporo, Japan). Briefly, the manufacturing process of FBRA is as follows. Fermentation base was made by steaming of brown rice and rice bran. Aspergillus Oryzae was then seeded to the fermentation base and fermentation process was continued for 18-24 h. Subsequently, second fermentation was continued for additional 12-24 h for aging purpose. Fermented product was then dried and powdered. 4-(Methylamino)-1-(3-pyridyl)-1-butanone (NNK) from Toronto Research Chemicals (Toronto, Canada), monoclonal rat anti-mouse Ki-67 antigen (Clone TEC-3), polyclonal rabbit anti-rat immunoglobulins/ biotinylated, polyclonal goat anti-rabbit immunoglobulins/ biotinylated, and Dako liquid DAB substrate-chromogen system were purchased from DAKO Corp. A polyclonal affinity-purified rabbit antibody to the cleaved caspase-3 (Asp175) was obtained from Cell Signaling Technologies. Vectastain® Elite ABC Kit was purchased from Vector Laboratories.

Experimental procedure. The experimental design was approved by the Institutional Ethics Review Committee for animal experiments at the Gifu University. Six weeks of age of female A/J mice were randomly divided into 8 groups as shown in Fig. 1. Mice in groups 2 and 3 received 5 and 10% FBRA in diet, respectively, during initiation stage (during and until 1 week after NNK treatment) and then transferred to the control diet until 21 weeks of age. Mice in groups 4 and 5, which were maintained on basal diet during NNK treatment, received 5 and 10% FBRA in diet, respectively, during the post-initiation stage (1 week after NNK treatment until 21 weeks of age). Mice in groups 7 and 8 received 5 and 10% FBRA in diet, respectively, throughout the experimental period. Mice in group 6 were maintained in control diet throughout the experimental period and served as non-treatment control. At 7 weeks of age, mice in groups 1-5 received the NNK (10 mmol) by i.p. injection. Fourteen weeks after NNK treatment, all surviving mice were sacrificed. Their lung were collected and weighted, infused with 10% neutral buffered formalin, and carefully inspected grossly. All of the macroscopically detected lung nodules were counted and examined histopathologically. Lung lesions, hyperplasias, and adenomas were diagnosed according to the criteria of ‘International Classification of Rodent Tumors: The Mouse’ (28). For Cyp2a5 level study, a total 10 female A/J mice were divided into 2 groups and received control diet and 10% FBRA in diet, respectively, at 6 weeks of age. Two weeks after FBRA treatment, all mice were sacrificed. Lung and liver tissues were collected for RNA isolation and quantitative analysis of Cyp2a5 expression.

RNA isolation and cDNA synthesis. Lung and liver tissue were cut off into small pieces (~50-100 mg) and extracted the RNA by using TRIzol reagent (Invitrogen, Life Technologies, Carlsbad, CA, USA) and an RNeasy Mini Kit (Qiagen Corp., Hilden, Germany) according to the manufacturer’s instructions. The amount and quality of RNA for each sample were assessed with the NanoDrop® ND-1000 UV-Vis Spectrophotometer (NanoDrop Technologies, Wilmington, DE, USA). Purified total RNA was reverse- transcribed using the SuperScript First-Strand Synthesis System for the reverse transcriptase-polymerase chain reaction (RT-PCR) kit (Invitrogen), according to the manufacturer's protocol.
Real-time quantitative PCR (RT-PCR). Expression levels of CYP2A5 and β-actin mRNA were measured using a LightCycler (Roche Applied Science) and the specific sense and antisense primers. The reaction mixture contained 10.0 μl of Takara SYBR Premix Ex Taq (Takara Bio Inc.), 1.0 μl of 10 μM of each primer, 3.0 μl of distilled water and 5.0 μl cDNA. The amplification number of cycles was 45 and the reaction took place for 10 sec at 95˚C, 10 sec at 60˚C and 6 sec at 72˚C, with an initial step of 95˚C for 10 min (hotstart). Oligonucleotide primers were the following: for CYP2A5, 5’-TGGTCCTGTATTCACCATCTACC-3’ and 5’-ACTACGCCATAGCCTTTGAAAA-3’; for β-actin, 5’-CATCCGTAGACCTCTATGCCAAC-3’ and 5’-ATGGAGCCACCGATCCACA-3’.

Immunohistochemistry. The avidin-biotin peroxidase complex (ABC) technique was used for immunohistochemical studies. The lung whitish nodules (size ≥0.8mm) from groups 1 and 5 were randomly collected. Sections (5-μm thick) were cut, deparaffinized, rehydrated in PBS, placed in 10 mmol/l citrate buffer (pH 6.0), and heated in Pascal pressure cooker (Code No.S2800, Dako Corp.) programmed for 1 min at 120˚C. The endogenous peroxidase activity was blocked by incubation for 20 min in 3% H2O2. After washing three times with PBS, the sections were preincubated with a normal blocking serum for 30 min at room temperature and then incubated with primary antibodies, which specific for Ki67 or cleaved caspase-3 (used at 1:100 dilution), overnight at 4˚C. Subsequently, the sections were incubated with biotinylated secondary antibodies (used at 1:250 dilution) for 30 min at room temperature, followed by incubation with avidin-coupled peroxidase (Vectastain Elite ABC Kit) for 30 min at room temperature. The sections were developed with 3,3V-diaminobenzidine (DAB), counterstained with hematoxylin and mounted coverslips.

Statistical analysis. The data for liver, kidney, lung or final body weights were analyzed by the Student’s t-test. The incidence, multiplicity and size of lung proliferative lesions were analyzed by Student’s t-test. Mouse CYP2A5 mRNA levels was analyzed by Mann-Whitney U-test. The Ki67 positive cell index was analyzed by Welch’s t-test.

Results

Final body weight and weight of the liver, kidney and lung. During the 21 weeks of experimental period starting at week 6, administration of FBRA caused no significant differences in final body weight and the weight of the liver, kidney or lung as shown in Table I. In addition, there were no clear signs of toxicity in the liver and kidney of any mice examined by the histological analysis.

Incidence, multiplicity and tumor size. Whitish nodules in the lung were frequently detected in mice treated with NNK. Only 1 mouse in the NNK-untreated group (group 7) harbored a nodule in the lung, suggesting a spontaneous development of lung tumor. There was no evidence of the development of lung tumors in groups 6 and 8. Histological analysis revealed that all lung lesions are adenomas (Fig. 2). The incidences and multiplicities (number of tumor per mice) of lung adenomas are summarized in Table II. The treatment of 10% FBRA (group 3) but not 5% FBRA (group 2) during the initiation phase significantly reduced the multiplicity of lung tumor when compared with control diet group 1 (p<0.006). However, 10% FBRA had no effects for the incidence and tumor’s size in comparison with group 1 as shown in Table II. The treatment of 10% FBRA during the post-initiation phase also significantly decreased in both the multiplicity and tumor size in comparison with group 1 (p<0.04) but not incidence of lung tumor induced by NNK (Table II).
Expression levels of Cyp2a5 mRNA in liver and lung. To investigate the molecular mechanisms of the suppressing effect of FBRA against NNK-induced lung tumorigenesis in the initiation phase, the mRNA expression levels of Cyp2a5 in both liver and lung tissues was analyzed by quantitative RT-PCR. Data for relative-quantification of Cyp2a5 mRNA in liver and lung of FBRA-treated A/J mice are summarized in Fig. 3. Administration of 10% FBRA led to the decreased
expression levels of Cyp2a5 mRNA in the lung (0.48±0.25 in FBRA treated mice vs. 1.28±0.68 in the control mice, p<0.02) but not in the liver (1.51±0.74 in FBRA-treated mice vs. 1.27±0.33 in the control mice).

Cell proliferative and apoptotic indices in lung tumors. In this study, suppressive effect of FBRA was confirmed not only in the initiation phase, but also in the post-initiation phase. Thus, cell kinetic analysis for cell proliferative and apoptotic index of lung tumors was performed with the expression of Ki67 and cleaved caspase-3. Administration of 10% FBRA to NNK-treated mice significantly decreased Ki67 positive cell index in the lung adenomas when compared with the adenomas in the NNK-treated group with control diet alone (0.065±0.016 and 0.114±0.025 per field at x400 magnification, respectively, p<0.001). In contrast to the frequent expression of positive cells for the cleaved caspase-3 in the control section of the colon, positive cells were scarcely found in NNK-induced lung tumors from different groups, indicating that cellular apoptosis rarely occurs in such tumors (data not shown).

Discussion

It is widely recognized that cigarette smoke contains a number of carcinogens, tumor initiators, promoters, co-carcinogens and mutagens. The mutagenicity of many constituents in tobacco smoke has been demonstrated in a variety of prokaryotic and eukaryotic test systems. The carcinogenicity of some of these mutagens has been evaluated in animals; some of these compounds are already designated as human carcinogens. 4-(Methylnitrosamino)-1-(3-pyridyl)-1-butanone (NNK) is one of the major carcinogenic compounds in cigarette smoke. Several epidemiological studies have suggested that NNK is one of the important causative agents for human lung cancers (2-5).

It is assumed that dietary factors play an important role in the prevention of several types of cancer, including lung cancer. A number of reports have shown the chemopreventive effect of fermented brown rice and rice bran (FBRA) on several types of cancer (22,24-26). The data presented herein also demonstrate that the exposure of 10% FBRA in the diet during both initiation and post-initiation phase significantly suppressed occurrence of the total number of lung proliferative lesions of NNK-treated A/J mice. In this study, the administration of 10% FBRA in the post-initiation phase significantly decreased the mean size of tumor. This inhibitory action is consistent with the previous reports that FBRA inhibited the carcinogenesis of the colon, liver, stomach, bladder and esophagus (22-26).

It is known that absorbed NNK initially exist in the body as a procarcinogen, an inert form that requires activation to exert its full effects. There are three primary pathways with regard to NNK activation: i) carbonyl reduction, ii) pyridine N-oxidation and iii) α-hydroxylation (5,29,30). Cytochrome pigment 450 (CYP450) enzymes belonging to the CYP multigene family catalyze the α-hydroxylation of NNK in the
oxidative metabolism pathway (5,9,11). It is noteworthy that expression levels of Cyp2a5 were significantly decreased by the administration of FBRA in this study. It is possible that the inhibitory effect on the Cyp2a5 expression leads to a decreased production of the active form of NNKs such as α-hydroxyNNKs and α-hydroxyNNALs, and is associated with the suppressive effects of FBRA on the lung tumorigenesis at the initiation phase.

In general, hyperproliferation is suggested to be relevant to carcinogenesis of many organs (31-33). Indeed, carcinomas of the lung have the highest indices for Ki67 (34,35). Therefore, the control of cell proliferation in the target organs is regarded as an important strategy for chemoprevention. Presently, we showed that dietary exposure of FBRA during the post-initiation phase decreased the size of tumors. It is important that the number of Ki67 positive cells in lung adenomas was significantly decreased by the post-initiation treatment of FBRA. Similar results have been obtained in other studies with various chemopreventive agents (36,37) and FBRA (24). The suppression of proliferative activity in tumor cells might be one of the important mechanisms of the chemopreventive effect of FBRA at post-initiation phase.

FBRA is a processed food prepared by fermenting brown rice and rice bran with Aspergillus Oryzae. The nutritional and sanitary advantage of fermentation has been recognized, although the details are not well known. Fermented soybean paste or soy sauce, which prepared by fermenting with Aspergillus Oryzae, has been found to be more stable against lipid peroxidation than unfermented soybeans, because Aspergillus Oryzae-fermented soybean products contain several antioxidants like 6-hydroxydaidzein, 8-hydroxydaidzein and 8-hydroxyisoflavone more abundantly than unfermented soybeans (38). These 8-hydroxylflavones are reported to possess greater 1,1-diphenyl-2-picrylhydrazyl (DPPH) radical-scavenging activity and antiproliferative activity than the corresponding isoflavone analogs (38,39). Furthermore, such 8-hydroxylflavones and 6-hydroxydaidzein are reported to possess antimutagenic activity (39,40). Several epidemiological and preclinical studies suggested the chemopreventive effects of such fermented soybean products (41-43). Dihydroferulic acid and dihydroresinamic acid, which were isolated from unpolished rice vinegar (Kurosu), were suggested as the major constituents responsible for radical scavenging activity of Kurosu (44). These acids are produced in Kurosu through the process of the fermentation from ferulic acid and sinapic acid. Rice bran contains approximately 20% oil which contains several bioactive polyphenols including ferulic acid, protocatechuic acid, sinapic acid and vanillic acid (45). Thus, it is possible that FBRA has similar mechanistic aspects to the fermented soybeans or Kurosu.

In conclusion, we demonstrated that administration of fermented brown rice and rice bran (FBRA) during both the initiation and post-initiation phase inhibits lung tumorigenesis induced by NNK in female A/J mice, indicating that FBRA is a promising chemopreventive agent for human lung cancer. Our results suggest that a reduction of Cyp2a5 mRNA expression levels and a reduction of cell proliferative activity in tumor cells may be involved in such chemopreventive effects of FBRA.

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