Potential for Preventing Diabetes and Dementia by Consuming Unpolished Rice Blended with Black Unpolished Rice and Super-Hard Rice

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ABSTRACT

Type-2 diabetes, which may prompt Alzheimer’s disease, can be regulated by diet. We investigated the palatability and efficacy of consuming two diets rich in β-glucan or resistant starch by blending ordinary polished rice and polished waxy barley (5:5) or by blending ordinary unpolished rice, unpolished pigmented rice and polished super-hard rice (2:4:4), then subjecting the blends to high-pressure treatment before cooking. After subjects consumed these diets for 12 weeks, consumers of the former rice blend had significantly less body fat, lower blood glucose levels. The latter blend showed higher antioxidant capacity and inhibition against β-secretase than control, and the subjects, consuming the latter blend, showed not only lower blood glucose levels, but also a higher ratio of amyloid β 42/40 peptide than the subjects consuming the traditional rice diet. Long-term consumption of the latter blended rice diets might inhibit rises in postprandial blood glucose levels and impede amyloid β production.

Abbreviations: Ae Mutant: Amylose Extender Rice Mutant, Aβ: Amyloid β Peptide Protein, BACE1: β-Secretase, HPT: High-Pressure Treatment, KBH Rice: “Koshihikari” Unpolished Rice Blended with Black Unpolished Rice And Super-Hard Rice (Ratio 2:4:4), KWB Rice: “Koshihikari” Polished Rice Blended with Polished Waxy Barley (Ratio 5:5), BGL: Blood Glucose Level, AUC: Area Under Rising Limb of a Glucose Curve for Postprandial Blood Glucose, AUCi: Area Under The Rising Limb of an Insulin Curve, FBG: Fasting Blood Glucose Level, RS: Resistant Starch, BFP: Body Fat Percentage, LDL: Low-Density Lipoprotein, HDL: High-Density Lipoprotein

Introduction

According to WHO report, globally, an estimated 422 million adults were living with diabetes in 2014, and diabetes of all types can lead to complications in many parts of the body and increase the overall risk of dying prematurely [1]. Therefore, there is an urgent need to implement population-based interventions that prevent diabetes, enhance its early detection, and use lifestyle and
High-pressure treatment (HPT), a technological process that limits the negative effects of food preparation on hydro-soluble vitamins, is recognized as being very useful in preserving nutritional quality in foods [22,23]. Yamakura et al. [24] showed that subjecting rice to HPT before cooking results in more free amino acids and stickier cooked rice.

In previous studies, we developed a novel method for inhibiting abrupt postprandial blood glucose levels (BGL) in Sprague–Dawley rats by feeding them α-mutant rice (super-hard rice) soaked with functional food ingredients that had been subjected to HPT before preparation [25]. Furthermore, when we pre-prepared (with HPT) “Koshihikari” rice blended with black rice (HPT KO) at a ratio of 6:4 (using a randomized, single-blind, crossover-designed study), we found that BGL in subject at 90 and 120 min after ingesting the cooked HPT KO was significantly lower than BGL following ingestion of the cooked “Koshihikari” (polished) rice alone. In fact, the increase in amyloid β40 peptide in blood 120 min after eating HPT KO rice tended to be lower than amyloid β40 after ingesting the cooked “Koshihikari” rice [26].

Recent epidemiological studies have suggested a link between Alzheimer’s disease (AD), and type 2 diabetes mellitus associated with insulin resistance [27-29]. Diabetes is a lifestyle disease, and its prevention and treatment are extremely important in improving longevity and quality of life. Low glycemic index (GI) foods inhibit rapid increases in blood glucose and insulin secretion after meals. The β-amyloid precursor protein (APP) generates the amyloid β peptide (Aβ) via β- and γ- secretase in a highly regulated process. Because working memory is impaired by AD, the disease has spawned dynamic research investigating the influence of gamma-aminobutyric acid (GABA) on working memory performance in AD sufferers [30]. It is well-recognized that the prevalence of dementia is higher in diabetic patients than in non-diabetic subjects [31], and Tokutake et al. [32] showed that there is a molecular link between AD and insulin signaling.

Molecules in rice with antioxidant activity include phenolic acids, flavonoids, anthocyanins, proanthocyanidins, tocopherols, tocotrienols, γ-oryzanol, and phytic acid. Rice bran also contains a variety of functional substances, such as γ-oryzanol, fericulic acid, sterol, wax, ceramide, phytin, inositol, and protein [33]. In previous studies, black rice bran contained more oleic acid and anthocyanin and showed greater BACE1 inhibitory activity than “Koshihikari” (a traditional, premium polished rice). Jensen et al. [34] and Qureshi et al. [35] showed that an increased daily consumption of bran significantly reduced the risk of coronary heart disease and lowered total cholesterol, LDL cholesterol, and triglycerides in serum. Abe et al. [36] showed that rice components that included inositol hexaphosphate significantly inhibited Aβ production in neuroblastoma cells without causing cytotoxicity, suggesting that such foods might prevent AD.

Pigmented rice contains naturally occurring pigmented flavonoids known as anthocyanins. Positive health effects have been reported for the pigments in the bran layer of rice. For example, consumption of red and black rice by rabbits has been found to reduce their atherosclerotic plaque formation and increase antioxidants, ostensibly by increasing concentrations of high-density lipoprotein cholesterol and apolipoprotein A1 [37]. In previous studies, we found that a super-hard rice bread blended...
with black rice bran (SRBBB), which contained a high amount of resistant starch (RS), provided a strong inhibitory activity against β-secretase (BACE1) and acetylcholinesterase, even after heating. Furthermore, aged mice fed the SRBBB diet for 4 weeks showed lower amyloid β40 peptide in their blood than mice fed a commercial feed diet [38].

In this study, we sought to develop a novel food product that could reduce abrupt increases in postprandial blood glucose spikes after meals for the main purpose of the experimental study, and as a side effect, those of inhibit Aβ protein production after meals. We examined the effects of two blended rice blend diets and a traditional diet of standard polished rice on insulin sensitivity, concentration of LDL to HDL cholesterol ratio, and body fat percentage (BFP). Both blended diets were cooked after HPT pretreatment of rice grains. One blended diet was comprised of super-hard rice blended with black unpolished rice (HPT KWB rice) and the other comprised of a β-glucan-enriched rice blend diet (HPT KWB rice). We used a 12-wk supplementation period with 1.6g RS/d, and with 2.9g β-glucan/d, compared with placebo, in 36 healthy subjects (slightly higher BGL) and assessed the results statistically.

**Materials and Methods**

**Materials**

We used four rice varieties and a mutant barley variety as food material. The a-e mutant rice cultivar “Chikushi-kona 85” was cultivated in 2017 in an experimental field at Fukuoka Agriculture and Forestry Research Center, Japan [9]. The high-dietary-fiber mutant rice cultivar “Goami 2”, developed in Korea, was provided to us by Dr. Hae Chune Choi in 2017. We purchased the high quality, premium, polished rice “Koshihikari” and the black rice “Okunomurasaki” from a local market. The waxy mutant barley cultivar “Hanenamochi” was cultivated in 2017 in an experimental field at the Hokuriku Research Center of the Central Agriculture Research Center, NARO, Japan.

**Preparation of Unpolished Rice Flour Samples and Polished Barley Grain Sample**

We prepared unpolished rice flour with a cyclone mill (SFC-S1; Udy Corp., Fort Collins, USA) screened with 1 mm diameter pores. Polished barley grain sample was prepared from barley seeds to be 70% yield ratio by barley pearling system 3RSB-10 (Hoden kogyo, Darmstadt, Germany).

**Food Processing of Experimental Diets**

We blended “Koshihikari” (unpolished rice) with “Okunomurasaki” (black unpolished rice), “Chikushi-kona 85” (unpolished, super-hard rice), and “Goami 2” (polished super-hard rice) at a ratio of 2:4:2:2, which we called KBH rice. We mixed “Koshihikari” (polished rice) with “Hanenamochi” (β-glucan-rich, polished waxy barley) at a ratio of 5:5, which we called KWB rice. Both rice mixtures were treated in a high-pressure machine (Ishikawajima-Harima Heavy Industries Co., Ltd., Tokyo, Japan) at 200 MPa for 2 min and cooked with an Echigo Seika, Co., Ltd. The standard-diet rice “Koshihikari” (polished and aseptically cooked) was purchased from a local market. All measurements and analyses were conducted in triplicate after sample preparation or cooking.

**Moisture Content of Rice Flour**

We measured the moisture content of the brown rice and cooked brown rice using the oven-drying method. Brown rice samples (2g) were dried for 1 h at 135°C, whereas cooked rice samples (2g) were dried for 3 h at 135°C. Cooked rice samples were stored at −80°C followed by lyophilizing the sample in a freeze dryer (FD-1; Tokyo Rikakikai Co., Ltd., Tokyo, Japan). Rice flour was prepared using a cyclone mill (SFC-S1; Udy Corp., Fort Collins, USA) screened with 1 mm diameter pores.

**Measurement of Iodine Absorption Spectra**

We measured the iodine absorption spectrum of milled rice with a Shimadzu UV-1800 spectrophotometer and estimated its apparent amylose content (AAC) using the iodine colorimetric method (as described by Juliano) [39]. The iodine absorption spectrum was analyzed [40] from 200 to 900 nm using a square cell with an inner dimension of 1 × 1 cm. (The control was distilled water in the same-sized cell.) We measured absorbance at λmax (i.e., peak wavelength) in the visible light range of iodine-stained starch because the intensity of iodine staining in starch has a high correlation with the length of glucan chains, the molecular size of amylose, and super-long chains (SLCs), as well as providing data on λmax, Aλmax, the λmax/Aλmax ratio. From these values, we introduce the term “New λmax,” which we derived as New λmax = (73.307 × Aλmax + 0.111 × λmax – 73.016)/ (λmax of various rice starches − λmax of glutinous starch).

**Glucose Content**

To each flour sample, we added 1 mL of 60% ethanol to a tube with 0.1 g of brown rice flour and then rotated the tube at 20°C for 1 h to extract glucose. The solution was then centrifuged at 1500 × g for 15 min, and the resulting supernatant was collected as a sample solution. We determined glucose content of each sample solution using the nicotinamide adenine dinucleotide phosphate enzyme assay method, measured with a glucose assay kit (Roche, Darmstadt, Germany).

**Physical Properties of Boiled Rice Grains**

We measured physical properties of cooked rice grains, based on 10 g of rice grains per sample, using a My Boy System Tensimeter (Taketomo Electric Co., Tokyo, Japan), according to the method described by Okadome et al. [41] The bulk measurements were taken six times, from which we calculated the mean for each sample.

**Measurement of RS**

RS of freeze-dried rice samples was measured according to the official method established by AOAC, using an RS assay kit (Mega-
zyme Ltd., Wicklow, Ireland). Each sample (0.1 g) was digested with pancreatin and amyloglucosidase at 37 °C for 6 h. Then, we measured glucose content using a spectrophotometer set at 510 nm.

**Beta-Glucan Assay Method**

We measured β-glucan in freeze-dried rice samples according to the official method of AOAC, using a β-glucan assay kit (Megazyme Ltd., Wicklow, Ireland). We measured β-glucan content by adding 0.2 mL of 50% ethanol to each rice flour sample (0.1 g) and then adding 5.0 mL of sodium phosphate buffer (20 mM, pH 6.5), stirring the tubes on a vortex mixer, and then incubating the tubes in a boiling water bath for approximately 2 min. Then, we removed the tubes and vigorously stirred them on a vortex mixer, heated the tubes at 50 °C for 5 min, and added 0.2 mL of lichenase (10 U) to each tube. We then capped the tubes, stirred, and incubated them at 50 °C for 1 h. We then added aliquot of 5.0 mL of sodium acetate buffer (200 mM, pH 4.0) and stirred the tubes on a vortex mixer. The tube remained at room temperature for 5 min and then was centrifuged at 1000 × g for 10 min. The supernatant was removed, and 0.1 mL aliquots were transferred to three test tubes; to one we added 0.1 mL of sodium acetate buffer (50 mM, pH 4.0) (the reaction blank) and to the other two tubes (the reaction) we added 0.1 mL of β-glucosidase (0.2 U) in 50 mM acetate buffer (pH 4.0). We then incubated the tubes at 50 °C for 10 min, added GOPOD reagent 3.0 mL to each tube, and incubated them at 50 °C for 20 min. We measured glucose content with a spectrophotometer set at 510 nm.

**BACE1 Inhibitory Activity**

We determined BACE1 inhibitory activity of freeze-dried rice samples using a BACE1 activity detection kit (Fluorescent; Sigma-Aldrich Co. LLC. Tokyo, Japan). To obtain the measurements, we first extracted 0.1 g of rice flour with 0.5 mL of 10 mM acetate buffer (pH 5.0) solution (0.1% Triton and 0.05% CHAPS) at room temperature for 1 h. Then, we centrifuged the solution at 10000 × g for 15 min and used an adsorption filter (MILLEX-GN33 mm: Merck Millipore Co Ltd. Tokyo, Japan) to remove the colored component from the extracted solution. We then added the sample to a 96-well tissue culture plate (AS ONE Corporation Osaka, Japan) to measure fluorescence. The fluorescent buffer (78 μL), extraction solution (30 μL) (the blank was set without a fluorescent buffer), and 0.03 U/μL enzyme (10 μL) were added to the samples (in this order) and then pre-incubated at 37 °C for 10 min. Next, 50 μM substrate (20 μL) was added to samples; samples were covered with Parafilm and then incubated at 37 °C for 2 h. We measured absorbance values at 320 nm (excitation) and 405 nm (emission) using a fluorescent microplate reader (Grating Based Multimode Reader SH-9000: CORONA ELECTRIC Co, Ltd. Ibaragi, Japan). The Trolox<sup>®</sup> calibration solutions, blank (dilute buffer), and diluted test samples (35 μL, fluorescein solution (115 μL, 110.7 nmol/L), and AAPH [2,2′-azobis (2-aminopropane) dihydrochloride] solution (50 μL, 31.7 mmol/L: H-ORAC, 63.4 mmol/L: L-ORAC). Measurements were performed immediately at 37 °C for 90 min.

**Measurement of Plasma Concentrations of Aβ1-42 And Aβ1-40**

Plasma concentrations of Aβ1-42 and Aβ1-40 were analyzed using V-PLEX Aβ Peptide Panel 1 (6E10) Kit (Meso Scale Discovery, Rockville, MD, USA) with MESO QuickPlex SQ 120 (Meso Scale Diagnostics, LLC, Rockville, MD, USA) according to the manufacturer's instruction. Intra-assay and inter-assay coefficients of variation were less than 20% for all assays.

**Study Design**

The study protocol for human test subjects was approved by the ethics committee for human tests of the Niigata Bio-research Park and Niigata University of Pharmacy and Applied Life Sciences and according to the 2014 guidelines issued by the Ministry of Education, Culture, Sports, Science, and Technology and the Ministry of Health, Labor, and Welfare, Japan. The test is registered as UMIN000030710. We recruited 36 normal, healthy human subjects.
(13 male, 23 female) for the study: fasting blood sugar level 78–119 mg/dL (mean = 101.94 mg/dL) and body mass index 15.5–27.3 kg/m² (mean = 21.4 kg/m²), grouped into three blocks. Subjects of each block ate the assigned test sample meals containing 50 g of carbohydrates once every day for 12 weeks consecutively. Subjects chose any other meals for two meals every day. After 0, 6, 12 weeks, test subjects got the single-dose test for BGL measurements, where they consumed the assigned meals within 10 min with frequent mastication (30 times was the guideline) and 200 mL of water. BGL and insulin levels were measured at 0, 30, 60, 90, and 120 min after three groups of 12 subjects had eaten three different test meals. Blood samples were acquired (in February and March of 2018) from each subject for analysis of Aβ40, Aβ42, HDL cholesterol, LDL cholesterol, and insulin sensitivity. BFP was measured with a body composition meter (DC-320 separate type, TANITA Co., Ltd., Tokyo, Japan) using the impedance method.

Statistical Analyses

All results were subjected to t-tests and Dunnett’s test using Excel Statistics (ver. 6, Microsoft Corporation, Tokyo, Japan). A value of p< 0.1 was considered to show the tendency, and a value of p< 0.05 was considered to be statistically significant.

Results and Discussion

Nutrient Intake and Structural Components of Test Meals

Table 1 β-glucan can reduce serum concentrations of total and LDL cholesterol [46] but the mechanism behind this reduction is unclear. However, it is difficult to predict the cholesterol-lowering effect of any particular food product enriched with β-glucan. [46] The Food and Drug Administration (FDA) of USA has stated that daily intakes of 3 g or more of soluble fiber (β-glucan) in barley may reduce the risk of heart disease due to its ability to lower total and LDL cholesterol. [47] Aoe et al. showed that the intake of high β-glucan barley (4.4 g/d) leads to significant and safe reductions in visceral fat area (VFA), body weight, BMI, and waist circumference (WC) for 12-wk. [46] β-glucan content was 9.7-fold higher in cooked HPT KB rice diet (2.9 g/d) than in the cooked control rice diet and the HPT KB cooked rice diet (0.3 g/d) diets (Table 1). β-glucan content and dietary fiber showed a similar tendency in response to diet treatment effects. In contrast, lipid content was 4.4-fold higher in cooked HPT KBH rice diet (2.2 g/d) than in the control diet (0.5 g/d). In addition, the HPT KBH rice diet showed high anthocyanin values (0.2 g/d).

![Table 1](image_url)

**Table 1**: Nutrient intakes and structural components of test-meals.

|                        | Control | Test Meal-1 | Test Meal-2 |
|------------------------|---------|-------------|-------------|
|                        | Koshihikari Polished Cooked Rice | HPT KBH Cooked Rice | HPT KWB Cooked Rice |
| Dietary intake (g)     | 151.0   | 139.0       | 168.0       |
| Carbohydrate (g)       | 50.0    | 50.0        | 50.0        |
| Dietary fiber (g)      | 0.5     | 2.1         | 4.6         |
| β-Glucan (g)           | 0.3     | 0.3         | 2.9         |
| Protein (g)            | 2.9     | 4.2         | 4.5         |
| Lipid (g)              | 0.5     | 2.2         | 0.8         |
| Sodium (mg)            | 6.8     | 6.9         | 5.0         |
| Anthocyanin (g)        | 0.0     | 0.2         | 0.0         |
| Moisture (g)           | 98.0    | 83.0        | 112.0       |
| Calorific value        | 228.0   | 225.0       | 227.0       |

Dietary fiber: enzymatic-gravimetric method (Proskyvariant)
β-glucan: Official method of AOAC using an β-glucan assay kit (Megazyme Ltd., Wicklow, Ireland).
Protein: Kjeldahl method.
Lipid: Gas chromatography.
Sodium: Inductively coupled plasma atomic emission spectrometry.
Anthocyanin: HPLC method.
Moisture: Drying method by heating.
Calorific value: (kcal/100g)= 4kcal × Protein + 9kcal × Lipid + 4kcal × Carbohydrate + 2kcal × Dietary fiber
Test meal-1: “Koshihikari” unpolished rice, black unpolished rice “Okunomurasaki”, and unpolished super-hard rice “Chikushi-kona 85”, and polished super-hard rice “Goami 2” blended at a ratio of 2:4:2 and cooked after high-pressure-treatment (HPT).
Test meal-2: HPT KB rice: polished rice “Koshihikari”, β-glucaSn-rich waxy hull-less barley, “Haneumamochi” blended in a ratio of 5:5 and cooked after a high-pressure-treatment (HPT).
Control meal: “Koshihikari” polished cooked rice.
Changes in Physical Findings and Blood Inspection Values of Three Groups of Test Subjects

Murao et al. [48]. showed that the LDL-to-HDL cholesterol ratio was 2.24 in subjects not being treated for diabetes mellitus, hypertension, or dyslipidemia, and that the ratio of subjects with no traditional cardiovascular risk factors (BMI, blood pressure, fasting blood glucose, HbA1c, HDL, LDL cholesterol, triglyceride, and uric acid levels below ideal levels) was 1.46. In comparison, after 12 weeks of test subjects following their assigned diets in our study, we determined that test subjects had LDL-to-HDL cholesterol ratio of 1.9 [SD (standard deviation): 0.8] for control diet subjects, 2.0 [SD: 0.5] for HPT KBH diet subjects, and 2.0 [SD: 0.8] for HPT KWB diet subjects (Table 2). Table 2 summarizes the effects of the intervention on the measured anthropometric and serum biochemical parameters. During the trial, body weight and BMI did not show significant difference among the three groups. As a result, the HPT KBH and HPT KWB diets had no effect on total or LDL cholesterol after 12 weeks following the diet.

Table 2: Changes in physical parameters and blood inspection values of the three test diets Changes in physical findings and blood inspection value of the test diet three groups of test subjects.

| Parameter                      | Test diet group | Baseline | After 6-wk | After 12-wk |
|-------------------------------|----------------|----------|------------|-------------|
| Body fat (%)                  | Control meal   | 30.0 ± 7.4 | 29.2 ± 8.2 | 30.2 ± 7.9  |
|                               | Test meal-1    | 25.8 ± 8.0 | 24.9 ± 8.1 | 25.3 ± 7.4  |
|                               | Test meal-2    | 26.2 ± 6.6 | 24.3 ± 6.6 | 24.8 ± 6.3  |
| Body weight (kg)              | Control meal   | 58.0 ± 5.8 | 57.7 ± 5.8 | 58.3 ± 6.2  |
|                               | Test meal-1    | 54.9 ± 8.6 | 54.7 ± 8.9 | 55.1 ± 9.2  |
|                               | Test meal-2    | 58.3 ± 12.3| 57.5 ± 11.6| 58.0 ± 11.6 |
| BMI (Body mass index) (kg/m²) | Control meal   | 22.1 ± 2.1 | 22.0 ± 2.10| 22.3 ± 2.2  |
|                               | Test meal-1    | 20.5 ± 3.4 | 20.4 ± 3.6 | 20.5 ± 3.7  |
|                               | Test meal-2    | 21.3 ± 2.8 | 20.9 ± 2.7 | 21.1 ± 2.7  |
| Pulse bpm, beats per min      | Control meal   | 67.5 ± 10.9| 69.5 ± 12.6| 64.4 ± 7.5  |
|                               | Test meal-1    | 67.7 ± 9.3 | 69.9 ± 11.5| 63.6 ± 7.2  |
|                               | Test meal-2    | 66.7 ± 11.0| 70.2 ± 9.0 | 65.1 ± 11.1 |
| Systolic blood pressure (mmHg)| Control meal   | 123.0 ± 15.2| 121.3 ± 16.1| 118.6 ± 13.7|
|                               | Test meal-1    | 120.0 ± 13.6| 119.7 ± 13.9| 111.9 ± 11.8|
|                               | Test meal-2    | 121.3 ± 17.4| 118.0 ± 15.4| 113.8 ± 12.8|
| Diastolic blood pressure (mmHg)| Control meal  | 74.9 ± 11.8 | 74.0 ± 13.2 | 72.1 ± 12.3  |
|                               | Test meal-1    | 71.5 ± 11.6 | 72.0 ± 9.8 | 67.0 ± 11.5  |
|                               | Test meal-2    | 73.3 ± 14.6 | 74.1 ± 10.5 | 67.5 ± 9.3  |
| Triglyceride [mg/dL]          | Control meal   | 718 ± 36.56 | 764 ± 50.0 | 78.6 ± 45.6  |
|                               | Test meal-1    | 69.1 ± 19.3 | 73.0 ± 23.4 | 81.3 ± 23.3  |
|                               | Test meal-2    | 71.8 ± 29.0 | 72.8 ± 30.0 | 82.3 ± 23.5  |
| LDL-Cholesterol [mg/dL]       | Control meal   | 718 ± 29.0 | 128.3 ± 38.2| 131.0 ± 38.9 |
|                               | Test meal-1    | 107.2 ± 20.7| 111.6 ± 19.6| 112.6 ± 19.1 |
|                               | Test meal-2    | 106.5 ± 21.0| 109.7 ± 25.4| 117.0 ± 29.2 |
| HDL-Cholesterol [mg/dL]       | Control meal   | 745 ± 15.8 | 74.6 ± 15.0 | 71.9 ± 14.9  |
|                               | Test meal-1    | 617 ± 10.3 | 599 ± 12.0 | 57.4 ± 11.2*  |
|                               | Test meal-2    | 67.2 ± 15.3 | 64.4 ± 15.0 | 63.9 ± 13.8  |
| Urea nitrogen [mg/dL]         | Control meal   | 1525 ± 2.4 | 127.2 ± 2.0 | 124.3 ± 3.6  |
|                               | Test meal-1    | 128 ± 2.9 | 127.2 ± 3.8 | 122.2 ± 2.8  |
|                               | Test meal-2    | 11.5 ± 1.6 | 13.6 ± 3.1 | 12.6 ± 2.6  |
| Total protein [g/dL]          | Control meal   | 7.3 ± 0.5 | 7.4 ± 0.3 | 7.3 ± 0.3  |
|                               | Test meal-1    | 7.3 ± 0.3 | 7.3 ± 0.3 | 7.1 ± 0.3  |
|                               | Test meal-2    | 7.0 ± 0.2 | 7.1 ± 0.2 | 7.2 ± 0.3  |

Control meal (3 male: 9 female), test meal-1 (4 male: 8 female), and test meal-2 (6 male: 6 female); significant by the Dunnett’s test (p<0.05).
Physical Properties of Cooked Rice

Cooked grains of ae-mutant rice cultivars are hard and non-sticky because they lack the starch-branching enzyme Iib and contain more SLCs. Ae-mutant grains appear to provide promising health benefits, such as diabetes prevention [7,8]. The physical properties of cooked rice, measured in bulk using a Tensipresser, detected differences among rice varieties in hardness, toughness, degree of adhesion, and stickiness, all important indices when evaluating rice palatability [41], with toughness showing similar tendencies as hardness. “Chilushi-kona 85” rice was the hardest and toughest among all samples tested, but it was also the least sticky and least adhesive (Table 3a). As a result, cooked grains of the ae-mutant brown rice cultivar are too hard and non-sticky to be marketed as table rice. Moreover, the hardness and toughness of the unpolished rice “Chilushi-kona 85” and polished rice “Goami2” were reduced 0.9-fold after HPT, whereas the adhesion and stickiness of black, unpolished rice were increased 3.3-fold and 2.4-fold after HPT, respectively (relative to the control). Furthermore, the hardness, toughness, adhesiveness, and stickiness for the waxy barley cultivar “Haneumamochi” increased by 1.2-fold after HPT relative to the untreated control (data not shown). We show that HPT treatment should make ae-mutant unpolished rice and black unpolished rice acceptable to consumers as table rice, after cooking. Therefore, HPT would be very beneficial to the rice industry due to the physical improvements HPT affords unpolished rice.

The hardness value of kernels was 4.8-fold higher in cooked HPT KBH rice than cooked rice of the control diet (“Koshihikari” polished rice). Toughness values were positively related to hardness, whereas adhesion was positively related to stickiness (Table 3b). However, the stickiness was 0.3-fold lower in cooked HPT KBH and KWP KBH rice than in cooked rice of the control diet. The polished waxy barley contained starch that was 97%–100% amylopectin; [49], for that reason, the hardness and toughness for HPT KBW rice were similar to the control rice diet.

Table 3b: Physical properties of individual material rice and blend rice by HPT and untreated cooked rice.

Table 3a:

|                    | Hardness [gw/cm²] | SD | Toughness [gw/cm²] | SD | Adhesion [gw/cm²] | SD | Stickiness [gw/cm²] | SD |
|--------------------|-------------------|----|--------------------|----|-------------------|----|---------------------|----|
| Koshihikari brown rice | 13.4              | a  | 22.9               | a  | 43.1              | a  | 17.1               | a  |
| Okunomurasaki brown rice | 25               | b  | 28.3               | a  | 6.5               | b  | 1.5                | b  |
| Chikushi-kona85’ brown rice | 96.6          | c  | 38.3               | b  | 0.3               | d  | 0.1                | c  |
| Goami2 white rice  | 14.3              | a  | 19.3               | a  | 2.7               | b  | 0.4                | b  |
| Koshihikari brown rice (HPT) | 16.6         | a  | 26.5               | a  | 8.9               | b  | 9.2                | b  |
| Okunomurasaki brown rice (HPT) | 22.2         | b  | 24                 | a  | 21.4              | c  | 10.2               | a  |
| Chilushi-kona 85’ brown rice (HPT) | 86.9        | c  | 34.5               | b  | 0.2               | d  | 0.1                | c  |
| Goami2 white rice (HPT) | 12.9              | a  | 16.3               | a  | 3.4               | b  | 2.4                | b  |

Means with the same letter are not significantly different (p<0.05). Control meal: “Koshihikari” polished cooked rice.

Measurement of Iodine Absorption Spectra

Starch properties measured by the iodine colorimetric method is shown in Table 4. AAC is an effective parameter for estimating the cooking or eating quality of rice grains. Because ae-mutants lack the starch-branching enzyme Iib, its amylopectin contains more long-chain glucans than does ordinary indica and japonica rice grains [50-53]. AAC concentration values for ae-rice cultivars “Chilushi-kona 85” (28.5%) and “Goami2” (35.1%) were higher than those of typical japonica rice cultivar “Koshihikari” (12.1%) and black rice “Okunomurasaki” (12.1%). The high molecular weight amylopectins in ae-mutant cultivars tend to have a higher value for λmax, Δλmax, New λmax, and Fb3 (the proportion of amylopectin with longer...
chains) (DP ≥ 37) [40], indicating that rice with high λmax/Åλmax ratios are palatable [40]. The λmax/Åλmax ratios for “Chikushi-kona 85” (1082.1) and “Goami2” (1049.9) were lower than ratios for “Koshihikari” (2189.3) and “Okunomurasaki” (2257.7). Furthermore, AAC concentration values for “Chikushi-kona 85” (18.3%) (pretreated with HPT) was 60% that of the control (28.5%) rice. The AAC values for the other samples (pretreated with HTP) showed almost the same values as the control samples.

Furthermore, the Aλmax, New λmax, and Fb3 (DP ≥ 37) values for “Chikushi-kona 85” (pretreated with HTP) values were 80% of the control. The Aλmax, New λmax, and Fb3 (DP ≥ 37) values for other samples (pretreated with HTP) showed almost the same values as the control samples, whereas the λmax/Åλmax ratios for all samples (pretreated with HTP) were 1.02–1.26 times higher than control samples. It appears from these data that pretreating rice with HPP renders the starch component of rice grains more palatable.

Table 4:
Analyses of iodine absorption curve and glucose content of individual material rice by HPT and untreated rice.

Table 4a:

| Material Rice          | AAC (%) | SD | λmax (nm) | SD | Åλmax | SD | λmax/Åλmax | SD | New λmax | SD | Fb3 (DP ≥ 37) (%) | SD |
|------------------------|---------|----|-----------|----|--------|----|------------|----|----------|----|-----------------|----|
| Koshihikari brown rice | 12.1    | a  | 561.5     | a  | 0.7    | a  | 0.257      | a  | 2189.3   | a  | 0.21            | a  |
| Okunomurasaki brown rice | 12.1    | a  | 565.5     | a  | 0.7    | a  | 0.251      | a  | 2257.7   | a  | 0.19            | a  |
| Chikushi-kona 85’ brown rice | 28.5    | b  | 567       | a  | 1.4    | b  | 0.524      | b  | 0.006    | b  | 0.65            | b  |
| Goami2 white rice      | 35.1    | c  | 579       | b  | 1.4    | b  | 0.552      | b  | 0.006    | b  | 0.57            | b  |
| Koshihikari brown rice (HPT) | 11.1    | a  | 559       | a  | 1.4    | a  | 0.245      | a  | 2286.4   | a  | 0.19            | a  |
| Okunomurasaki brown rice (HPT) | 12    | a  | 570.5     | b  | 3.5    | b  | 0.247      | a  | 2309.7   | a  | 0.11            | a  |
| Chikushi-kona 85’ brown rice (HPT) | 18.3    | d  | 557.5     | a  | 2.1    | c  | 0.41       | c  | 0.001    | c  | 0.53            | b  |
| Goami2 white rice (HPT) | 34.5    | c  | 581.5     | d  | 2.1    | b  | 0.537      | b  | 1082.9   | b  | 1.1             | b  |

Table 4b:

| Glucose Content (G/100g) | SD       |
|--------------------------|----------|
| Koshihikari brown rice (HPT) | 0.03     |
| Okunomurasaki brown rice (HPT) | 0.07     |
| Chikushi-kona 85’ brown rice (HPT) | 0.18     |
| Goami2 white rice (HPT) | 0.284    |
| Haneumamochi | 0.23     |

Changes in Postprandial BGL, Insulin Levels, Area Under the Curve (AUC), And the Insulin Rise Curve (AUCi) In Humans

Panlasigui and Thompson [54] showed that unpolished rice is healthier and more beneficial than polished rice for diabetics and hyperglycemic individuals and that the glycemic area and GI were 35.4% lower in unpolished rice than in polished rice. Jung et al. [55] showed that ferulic acid might be beneficial in the treatment of type-2 diabetes because it regulates BGLs by elevating glucokinase activity and the production of glycogen in the liver. The BGL of test subjects 30 min after ingesting cooked HPT KBH (test meal-1) showed lower tendency after 12 weeks on the diets than that in subjects who consumed the control diet (p< 0.1), and that of cooked HPT KWB (test meal-2) was significantly lower than that of control diet (p< 0.01) (Figure 1A), whereas AUC showed a similar tendency in response to diet after 30 and 60 min (Figure 1B). After 12 weeks on treatment diets, insulin levels of test subjects that ingested cooked HPT KBH rice (test meal 1) were significantly lower at 120 min after the meal than insulin levels in subjects consuming the control diet (p < 0.05) (Figure 2A). Furthermore, after 12 weeks on their assigned diets, AUCi in test subjects at 90 and 120 min after ingesting the cooked HPT KWB (test meal 2) showed lower tendency than those in test subjects who ingested the control diet (p < 0.1, Figure 2B).
RS Contents of Samples

RS is starch that eludes digestion in the small intestine and may ferment in the large intestine. Several studies have reported that long-term consumption of RS might reduce fasting cholesterol and triglyceride concentrations. [19-21] Reader et al. [56] showed that 7.25 g/d of RS added to an energy bar reduced blood glucose and insulin levels in healthy adults. The RS contents of the “Goami2” (9.8%) and “Chikushi-kona 85” (10.2%) were very high (i.e., 44.3-46.3-fold higher than RS in “Koshihikari” unpolished rice (0.2 %) (Table 5). RS contents for “Koshihikari” unpolished rice, “Chikushi-kona 85” unpolished rice, and “Haneumamochi” were 1.5-fold and 1.2-fold higher (relative to the control), respectively, after HPT (200 MPa for 2 min). RS content was 2.4-fold higher (P < 0.05) in cooked HPT KBH rice (2.9%) than in the cooked control rice (2.0%) (Table 7). In contrast, RS content was 0.8-fold lower in the cooked HPT KWB rice than in the cooked control rice diet.
Table 5: Resistant starch of individual material rice by HPT and untreated cooked rice.

| Material           | Resistant Starch (%) | SD |
|--------------------|----------------------|----|
| Koshihikari brown rice | 0.2                  | a 0|
| Okunomurasaki brown rice | 0.3                  | a 1|
| Chikushi-kona 85' brown rice | 10.2                 | b 0.5|
| Goami2 white rice    | 9.8                  | b 0|
| Haneumamochi        | 0.6                  | a 0|
| Koshihikari brown rice (HPT) | 0.3                  | a 0|
| Okunomurasaki brown rice (HPT) | 0.3                 | a 0|
| Chikushi-kona 85' brown rice (HPT) | 12                  | b 0.1|
| Goami2 white rice (HPT) | 9.9                  | b 0|
| Haneumamochi (HPT)   | 0.7                  | a 0|

Means with the same letter are not significantly different (p<0.05).

Control meal: “Koshihikari” polished cooked rice.

HPT KBH rice: unpolished rice “Koshihikari”, black unpolished rice, “Okunomurasaki”, un-polishesuper-hard rice, “Chikushkona85’ ” and polished super-hard rice, “Goami2.” blended in a ratio of 2:4:2:2 and cooked after a high-pressure-treatment (HPT).

HPT KWB rice: polished rice “Koshihikari”, β-glucan-rich waxy hull-less barley, “Haneumamochi” blended in a ratio of 5:5 and cooked after a high-pressure-treatment (HPT).

Changes in Variation of Amyloid β42/40 Peptide Ratio Species

Amyloid beta peptides (Aβ) are important components of plaque in AD. Plasma concentration of Aβ (1–40) and Aβ (1–42) rise with age and are higher in people with mutations that cause early onset of AD. However, Aβ (1–42) concentrations may decline early in the dementia process [59]. High plasma concentrations of Aβ (1–40), especially when combined with low concentrations of Aβ (1–42), indicate an increased risk of dementia [59, 60]. Thus, plasma Aβ 42/40 ratio may be a useful premorbid biomarker for identifying cognitive deterioration [61]. Blood samples were taken from the 24 subjects after meals before diets began (baseline) and after 6 and 12 weeks on their assigned diets. Variation in Aβ 42/40 ratio after 6 weeks of consuming the HPT KBH rice diet (0.0009) was significantly higher (p < 0.04) than ratio for subjects consuming the control diet (-0.0026) (Figure 4). Thus, changes in BACE1 inhibitory activity, due to consuming the HPT KBH rice diet over the long-term, might have played a role in lowering Aβ ratios after meals.

Measurement of Hydrophilic and Lipophilic Oxygen Radical Absorbance Capacity

Many dietary compounds have been proposed as being important antioxidants, and although there is credible evidence...
that vitamins E and C are important antioxidants, evidence is weaker for carotenoids and related plant pigments [62]. L-ORAC values of “Koshihikari” unpolished rice, “Okunomurasaki” unpolished rice, “Chikushi-kona 85” unpolished rice, and “Goami2” polished rice increased markedly after HPT (Table 6). Therefore, HPT increases lipophilic antioxidant substances, except in the waxy barley (“Haneumamochi”). This lack of increase may have been due to damage to cell membranes organelles, such as spherosomes, which liberate or create free lipophilic antioxidants, such as tocopherol. Conversely, H-ORAC values in the unpolished black rice “Okunomurasaki” were much higher than in other samples. It seems that anthocyanin in unpolished black rice plays a principal role in H-ORAC values because hydrophilic antioxidant anthocyanins react more effectively when subjected to HPT (200 MPa). L-ORAC for HPT KBH rice and HPT KWB rice increased 2.9-fold and 1.7-fold (P < 0.05), respectively, relative to the control rice diet (Table 7). H-ORAC values trended in similar directions to L-ORAC values in response to dietary treatment effects. Tsuda et al. [63] showed that anthocyanins extracted from purple corn prevented obesity and ameliorated hyperglycemia in mice. Moreover, Suzuki et al. [64], showed that the calcium content of pigmented brown rice was higher than the “Koshihikari” cultivar. These experiments suggest that anthocyanins may function in helping prevent oxidation in vivo.

**Table 6:** H-ORAC and L-ORAC capacity of individual material rice by HPT and untreated cooked rice.

|                     | H-ORAC μmol TE/100gFW | SD   | L-ORAC μmol TE/100gFW | SD   |
|---------------------|------------------------|------|------------------------|------|
| Koshihikari brown rice | 41.6                   | a 1.8 | 4.4                    | a 0  |
| Okunomurasaki brown rice | 285.8                  | a 8.6 | 5.3                    | a 0  |
| Chikushi-kona 85’ brown rice | 210.4                  | b 5.2 | 1.3                    | a 0  |
| Goami2 white rice            | 252.6                  | b 3.7 | 0.9                    | a 0  |
| Haneumamochi                     | 59.1                   | a 2.8 | 17.5                   | c 0.8|
| Koshihikari brown rice (HPT)    | 43.1                   | a 2.8 | 43                     | a 3.8|
| Okunomurasaki brown rice (HPT)  | 361                    | b 12.3 | 32.7                   | a 4  |
| Chikushi-kona 85’ brown rice (HPT) | 189                  | c 9.6 | 11.7                   | b 1.8|
| Goami2 white rice (HPT)               | 214.4                  | c 8.3 | 8.2                    | b 2.8|
| Haneumamochi (HPT)                | 50.5                   | a 3.8 | 11.1                   | b 3.1|

Means with the same letter are not significantly different (p<0.05).
Control meal: “Koshihikari” polished cooked rice.
HPT KBH rice: unpolished rice “Koshihikari”, black unpolished rice, “Okunomurasaki”, un-polish记者了解-hard rice, “Chikushkona85’ ” and polished super-hard rice, “Goami2” blended in a ratio of 2:4:2:2 and cooked after a high-pressure-treatment (HPT).
HPT KWB rice: polished rice “Koshihikari”, β-glucan-rich waxy hull-less barley, “Haneumamochi” were blended in a ratio of 5:5 and cooked after a high-pressure-treatment (HPT).

**Table 7:** Bio-functional properties of blended cooked samples.

|                     | Resistant starch (%) | SD | H-ORAC μmol TE/100gFW | SD | L-ORAC μmol TE/100gFW | SD | Glucose content (g/100g) | SD | β-Secretase inhibition rate (%) (43.4mg- eq./mL) | SD | H-ORAC μmol TE/100gFW | SD |
|---------------------|----------------------|----|------------------------|----|------------------------|----|-------------------------|----|-----------------------------------------------|----|------------------------|----|
| Control diet        | 1.2                  | a 0.1 | 9.3                    | a 0.8 | 0                      | a 0 | 0.14                    | a 0 | 0                                      | a 0 |
| HPT KWB             | 0.9                  | a 0   | 36.7                   | b 0.8 | 1.7                    | b 0 | 0.32                    | b 0.02 | 5                                      | b 0.7 |
| HPT KBH             | 2.9                  | b 0   | 77.2                   | c 0.8 | 2.9                    | c 0 | 0.2                     | a 0.01 | 19                                      | a 2.1 |

Means with the same letter are not significantly different (p<0.05).
Control meal: “Koshihikari” polished cooked rice.
HPT KBH rice: unpolished rice “Koshihikari”, black unpolished rice, “Okunomurasaki”, un-polish了解更多-hard rice, “Chikushkona85’ ” and polished super-hard rice, “Goami2” blended in a ratio of 2:4:2:2 and cooked after a high-pressure-treatment (HPT).
HPT KWB rice: polished rice “Koshihikari”, β-glucan-rich waxy hull-less barley, “Haneumamochi” blended in a ratio of 5:5 and cooked after a high-pressure-treatment (HPT).
Glucose Content and BACE1 Inhibitory Activity

Glucose concentrations of cooked rice samples were high in “Goami2” polished rice (0.284 g/L), “Haneumamochi” (0.230 g/L), and “Chikushi-kona 85” unpolished rice (0.180 g/L), whereas glucose content in “Koshihikari” unpolished rice (0.030 g/L) was very low (Table 4b). The glucose content remained almost identical between HPT (200 MPa 2 min) samples and untreated cooked rice (data not shown). However, glucose content was 2.3-fold higher (P < 0.05) in cooked HPT KWB rice (0.32 g/L) than in the cooked rice (0.14 g/L) of the control diet (Table 7). BACE1 is the first protease involved in the process of converting Aβ in the brain. BACE1 inhibitory activity for the HPT KBH rice (19.0%), HPT KWB rice (5.0%), and control diet (0.0%), indicated that the BACE1 inhibitory activity of pigmented rice bran is much stronger than inhibitory activities in ordinary rice bran, as reported in our previous paper [38] (Table 7).

Conclusion

Because it has been reported that type-2 diabetes increases the risk of AD, we investigated whether Aβ generation could be prevented by consuming foods resistant to digestion. To identify natural, neuroactive and slowly digestive compounds with a potent inhibitory effect on BACE1, we evaluated black unpolished rice, ae-mutant unpolished rice, and polished waxy barley as dietary alternatives to traditional polished rice and that it would prevent abrupt increases in postprandial BGL, insulin levels, and inhibitory activity against Aβ production, if consumed consistently as a regular, staple, component of one’s diet. In addition, HPT KBH cooked rice might help prevent dementia due to the inhibitory effects we found in the Aβ 42/40 ratio in test subjects that consumed the diet for 12 weeks. We will continue this investigation on multiple-preventive rice products against diabetes and dementia by the combination of black rice and barley in the near future.

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