

Effects of *Bacillus subtilis* var. natto products on symptoms caused by blood flow disturbance in female patients with lifestyle diseases

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Abstract: The fermented soybean product natto is a popular traditional food in Japan and is considered a health supplement. NKCP®, a natto-derived dietary food supplement whose main component is bacillopeptidase F, has antithrombotic, fibrinolytic, and blood pressure-lowering effects. We examined whether daily intake of NKCP® effectively improves subjective symptoms in patients with lifestyle diseases in this cross-over, double-blind study. Fermented soya extract with subtilisin NAT (nattokinase) as the main component was used as an active placebo. A 4-week course of NKCP® significantly decreased the visual analog scale (VAS) score for shoulder stiffness from 42.3 to 32.4 ($P=0.009$), the VAS score for low back pain from 25.5 to 18.8 ($P=0.02$), and the VAS score for coldness of the extremities from 33.1 to 25.7 ($P=0.002$). However, no significant difference was found in the VAS score for headache. After a 4-week course of active placebo, no significant changes in the VAS score were found for any symptoms. The significant improvement in the symptoms secondary to blood flow disturbance was caused by the improvement in blood flow by NKCP®. The use of dietary supplements based on the Japanese traditional food natto helps to relieve subjective symptoms for patients with lifestyle diseases receiving medical care.

Keywords: bacillopeptidase F, Japanese traditional food, lifestyle disease, subjective symptom, supplement

Introduction

Recently, traditional Japanese foods have attracted attention because of Japan's lower prevalence of cardiovascular diseases and the Japanese population's longer life expectancy that exceed life expectancy in Western countries.^{1,2} In 2013, the food culture of Japan was added to the United Nations Educational, Scientific, and Cultural Organization (UNESCO) List of Intangible Cultural Heritage.³ Natto, the fermented soybean product, is a popular traditional food in Japan. When forming natto, the soya beans are first soaked and cooked, then friendly starter bacteria are added, and the soya beans are allowed to develop under warm and damp conditions. Natto is considered a health supplement because of its high amount of nutrients: 100 g of natto contains a variety of amino acids (16 g), vitamin B₁ (0.07 mg), vitamin B₂ (0.56 mg), vitamin B₆ (0.24 mg), fibers (6.7 g), and so on. *Bacillus subtilis* var. natto, a bacterium isolate from natto, secretes several proteases during soybean fermentation. Of them, bacillopeptidase F and subtilisin NAT (nattokinase) are considered to have fibrinolytic activity.^{4–6} Both substances are supplied as dietary supplements because of their positive effects on blood flow; the former is present in NKCP®, and the latter is present in fermented soya extract.^{4–6} The anticoagulant, fibrinolytic, and blood pressure-lowering effects of both substances have

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been compared, and NKCP® has been scientifically confirmed to have dominant effects.^{7,8} NKCP® has a 2.5-fold greater fibrinolytic effect and a >100-fold anticoagulant effect in human blood; it also causes a significantly greater reduction in blood pressure in patients with lifestyle diseases including essential hypertension, type 2 diabetes, or hyperlipidemia than does fermented soya extract.^{7,8}

Because endothelial and platelet dysfunction as well as coagulation and fibrinolysis abnormalities more often occur in patients with than without hypertension, diabetes mellitus, or hyperlipidemia, we speculated that changes in blood fluidity improve symptoms caused by blood flow disturbances in such patients.⁹ In this study, therefore, we examined whether daily intake of the nutritional supplement NKCP® helps to improve subjective symptoms in patients with lifestyle diseases.

Methods

Study design

Patients who regularly visit the hospital or clinics with essential hypertension, type 2 diabetes, or hyperlipidemia were considered eligible for this study. This study was performed at three facilities in Japan (Nagareyama Central Hospital, Nagareyama; Ichihashi Clinic, Kobe; and Hanzoumon Clinic, Tokyo). Essential hypertension is defined as systolic blood pressure of 140 mmHg or more and/or diastolic blood pressure of 90 mmHg or more. Hyperlipidemia was defined as low-density lipoprotein cholesterol greater than 140 mg/dL,

triglycerides greater than 150 mg/dL, or high-density lipoprotein cholesterol less than 40 mg/mL. Type 2 diabetes was defined as fasting plasma glucose greater than 126 mg/dL, random plasma glucose greater than or equal to 200 mg/dL, or 2-hour glucose in the oral glucose tolerance test greater than or equal to 200 mg/dL. We excluded patients who were pregnant or possibly pregnant, were enrolled in another clinical trial, or patients with unstable angina, acute myocardial infarction, chronic obstructive pulmonary disease, renal insufficiency, acute liver disease, infectious disease, allergies to drugs or food, or any other serious life-threatening illness. Finally, 17 female patients with a mean age of 67.8 years (range, 55–85 years) participated in this study. The characteristics of all participants are shown in Table 1. All patients were nonsmokers and took concomitant medications, including lipid-lowering agents, antidiabetic drugs, or/and antihypertensive drugs. No changes were made to the pharmacological regimens of the patients during the course of the study.

Substance description

The substance used in the present study was NKCP® (250 mg/day, twice daily), the main component of which is bacillopeptidase F (Daiwa Pharmaceutical Co, Ltd, Tokyo, Japan). Fermented soya extract (100 mg/day, twice daily), the main component of which was subtilisin NAT, was used as the active placebo (Japan Bio Science Laboratory Co, Ltd, Osaka, Japan). Each dose was the quantity recommended

Table 1 Characteristics of participants

No	Age	Body height (cm)	Body weight (kg)	Symptom		
				Hypertension	Type 2 diabetes	Hyperlipidemia
1	76	150	47			○
2	69	150	48	○		
3	85	151	45	○		○
4	68	155	57	○		○
5	68	159	48	○		
6	61	160	71	○		○
7	71	153	52	○		○
8	63	148	56	○		○
9	59	150	66	○		
10	77	138	63	○	○	○
11	74	157	45	○	○	○
12	71	149	48	○		○
13	55	163	76	○	○	
14	62	152	62	○		
15	71	162	53	○	○	
16	55	154	49			○
17	68	162	76	○	○	

Abbreviations: No, identifying number of patient; O, suffering from this symptom.

by each manufacturer. The patients could not discriminate between the two substances. The cross-over, double-blind study was performed. The patient received one substance for 4 weeks, was observed for 4 weeks without receiving a substance, and then received another substance for 4 weeks. There were no adverse substance-related effects. Written informed consent was obtained from all patients before participating in the study. This study design was approved by the Shiba Palace Clinic Ethics Committee (Tokyo, Japan).

Parameters

Body height and weight were measured in the morning, and blood samples for laboratory assays were obtained before and after the administration of each substance. Samples were obtained following an overnight fast. Blood tests consisted of blood count (white blood cells, red blood cells, hematocrit, and hemoglobin); blood chemistry (total protein, albumin, blood urea nitrogen, creatinine, total bilirubin, aspartate aminotransferase, alanine aminotransferase, γ -glutamyl transpeptidase, lactate dehydrogenase, alkaline phosphatase, cholinesterase, creatine kinase, total cholesterol, low-density lipoprotein cholesterol, high-density lipoprotein cholesterol, triglycerides, blood sugar, and glycated hemoglobin [HbA_{1c}]); and D-dimer assay. Before and after administration of each substance, the patients reported their state of headache, shoulder stiffness, low back pain, and coldness of the extremities. A visual analog scale (VAS), anchored with “no symptom” on the left and “extreme symptom” on the right, was used to rate the intensity of the symptoms. The patients were asked to schematically plot their perceived symptom level at rest.

Statistical analysis

The Wilcoxon signed rank test was used to compare the laboratory and physical data or VAS scores before and after using each substance. Differences with a *P*-value of <0.05 were considered to be statistically significant.

Results

Physical findings

The mean height of all participants was 153.7 ± 1.5 cm (mean \pm standard error), mean weight was 56.5 ± 2.6 kg, mean body mass index was 23.9 ± 1.1 . When using each substance, mean body weight of the participants did not change significantly (NKCP® range, 56.6–56.8 kg, *P*=0.44; active placebo range, 56.5–56.4 kg, *P*=0.81). Regarding the blood pressure of the participants, a 4-week course of NKCP® significantly decreased the mean value of systolic blood pressure from 130.3 to 111.8 mmHg (*P*=0.002) and

that of diastolic blood pressure from 72.9 to 69.1 mmHg (*P*=0.024). However, after a 4-week course of active placebo, no significant changes in either systolic or diastolic blood pressure were found (124.1 vs 123.1 mmHg, and 70.1 vs 68.6 mmHg, respectively).

Blood tests

The average baseline complete blood count, protein level, liver function parameters, renal function parameters, and electrolyte level were normal before the administration of each substance (Table 2). The D-dimer levels and blood glucose levels before using NKCP® and active placebo, and the triglyceride levels before using active placebo, were higher than their respective reference ranges. Other values were within the reference ranges. No significant differences were found when the baseline values of all laboratory findings were compared between the NKCP® and active placebo group.

Regarding the comparison before and after using the substances, no significant differences were found in the laboratory parameters before and after the 4-week administration of NKCP®, with the exception of cholinesterase and HbA_{1c} (*P*=0.03 and 0.04, respectively). However, the values of these two parameters were within the reference ranges. Likewise, no significant differences in these parameters were found in patients taking active placebo, with the exception of the platelet count and cholinesterase level, which were within the reference ranges (*P*=0.01 and 0.02, respectively; Table 1).

Subjective symptoms

First, no significant differences were found in the baseline VAS scores for headache, shoulder stiffness, low back pain, or coldness of the extremities between NKCP® and active placebo group. Next, changes in the VAS scores were compared after the administration of NKCP® and active placebo (Figures 1–4). A 4-week course of NKCP® significantly decreased the VAS score of shoulder stiffness from 42.3 to 32.4 (*P*=0.009), that of low back pain from 25.5 to 18.8 (*P*=0.02), and that of coldness of the extremities from 33.1 to 25.7 (*P*=0.002). However, no significant difference was found in the VAS score for headache (*P*=0.17). For the active placebo group, no significant changes in the VAS score for each symptom were found after the 4-week course.

Discussion

With the increase of the aging in developed countries, chronic diseases, such as diabetes, hypertension, cardiovascular disease, and osteoporosis are also on the rise. Most of patients of these diseases suffer from any symptoms in accordance

Table 2 Changes in blood test parameters between baseline and after 4 weeks of administration of NKCP® and active placebo

Parameter	NKCP*			Active placebo		
	Baseline	4 weeks	P-value	Baseline	4 weeks	P-value
Red blood cell count ($\times 10,000 \mu\text{L}$)	440.9 \pm 11.2	437.4 \pm 11.3	0.168	435.1 \pm 11.9	442.1 \pm 11.6	0.155
White blood cell count ($/\mu\text{L}$)	5,717.1 \pm 28.22	5,963.5 \pm 217.6	0.436	5,685.9 \pm 253.7	5,768.2 \pm 284.5	0.667
Hemoglobin content (g/dL)	13.4 \pm 0.30	13.3 \pm 0.31	0.211	13.3 \pm 0.31	13.5 \pm 0.31	0.138
Hematocrit (%)	40.6 \pm 0.83	40.2 \pm 0.81	0.104	40.1 \pm 0.85	41.0 \pm 0.83	0.049*
Blood platelet count ($\times 10,000 \mu\text{L}$)	21.2 \pm 0.74	21.8 \pm 0.64	0.667	21.9 \pm 0.67	22.8 \pm 0.75	0.007**
Total protein (g/dL)	7.2 \pm 0.09	7.1 \pm 0.08	0.395	7.1 \pm 0.08	7.3 \pm 0.12	0.055
Albumin (g/dL)	4.4 \pm 0.06	4.3 \pm 0.07	0.333	4.3 \pm 0.05	4.4 \pm 0.08	0.053
Total bilirubin (mg/dL)	0.68 \pm 0.03	0.72 \pm 0.05	0.402	0.71 \pm 0.04	0.65 \pm 0.04	0.213
AST (U/L)	28.9 \pm 2.51	28.5 \pm 3.06	0.617	30.4 \pm 4.49	32.4 \pm 6.15	0.486
ALT (U/L)	25.1 \pm 3.90	24.6 \pm 4.14	0.445	25.6 \pm 4.76	26.2 \pm 5.26	0.434
ALP (U/L)	239.8 \pm 14.5	248.5 \pm 19.5	0.842	245.3 \pm 17.8	244.5 \pm 14.2	0.754
LDH (U/L)	212.8 \pm 8.4	205.5 \pm 6.6	0.129	205.8 \pm 8.1	206.8 \pm 7.9	0.947
γ -GTP (U/L)	31.1 \pm 5.28	32.3 \pm 5.25	0.578	32.0 \pm 5.70	34.4 \pm 6.36	0.953
Cholinesterase (U/L)	348.6 \pm 16.9	341.2 \pm 16.4	0.028*	340.1 \pm 17.5	355.2 \pm 20.2	0.022*
Creatine kinase (U/L)	138.8 \pm 21.3	119.9 \pm 14.1	0.358	127.4 \pm 17.9	125.5 \pm 16.5	0.509
Blood urea nitrogen (mg/dL)	15.6 \pm 0.91	16.0 \pm 1.05	0.895	15.3 \pm 0.88	15.2 \pm 0.77	0.965
Creatinine (mg/dL)	0.71 \pm 0.05	0.73 \pm 0.06	0.225	0.71 \pm 0.05	0.70 \pm 0.04	0.521
Total cholesterol (mg/dL)	204.6 \pm 7.5	200.1 \pm 6.3	0.161	198.6 \pm 5.2	197.3 \pm 6.9	0.598
HDL cholesterol (mg/dL)	58.8 \pm 2.50	56.6 \pm 2.65	0.333	58.1 \pm 2.60	60.5 \pm 2.69	0.336
LDL cholesterol (mg/dL)	117.9 \pm 6.1	113.7 \pm 5.9	0.088	114.5 \pm 5.4	112.1 \pm 5.9	0.559
Triglyceride (mg/dL)	148.4 \pm 16.6	181.9 \pm 33.6	0.493	153.9 \pm 27.0	138.0 \pm 15.8	0.582
Blood sugar (mg/dL)	109.6 \pm 11.0	112.2 \pm 9.2	0.982	116.9 \pm 12.6	105.6 \pm 7.8	0.768
Hemoglobin A _{1c} (%)	5.6 \pm 0.18	5.6 \pm 0.20	0.039*	5.6 \pm 0.21	5.6 \pm 0.20	0.366
D-dimer ($\mu\text{g/mL}$)	0.58 \pm 0.04	0.61 \pm 0.05	0.348	0.55 \pm 0.02	0.57 \pm 0.04	0.668

Notes: Values are represented as mean \pm standard error. Statistical significance, * $P<0.05$; ** $P<0.01$.

Abbreviations: AST, aspartate aminotransferase; ALT, alanine aminotransferase; ALP, alkaline phosphatase; LDH, lactate dehydrogenase; γ -GTP, gamma-glutamyl transpeptidase.

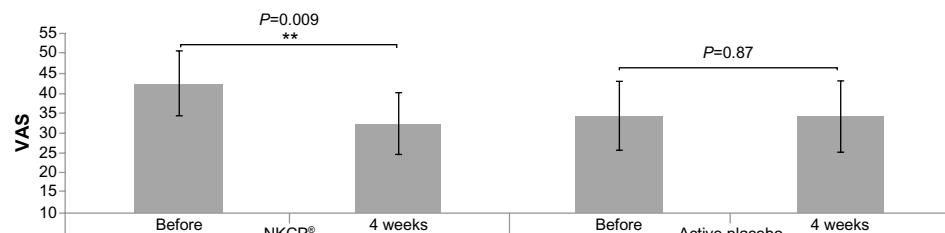


Figure 1 Changes in VAS score for shoulder stiffness after 4 weeks compared with baseline in patients taking NKCP® (Daiwa pharmaceutical Co., Ltd.) or active placebo.

Notes: The vertical bar indicates the standard error. Statistical significance, ** $P<0.01$.

Abbreviation: VAS, visual analog scale.

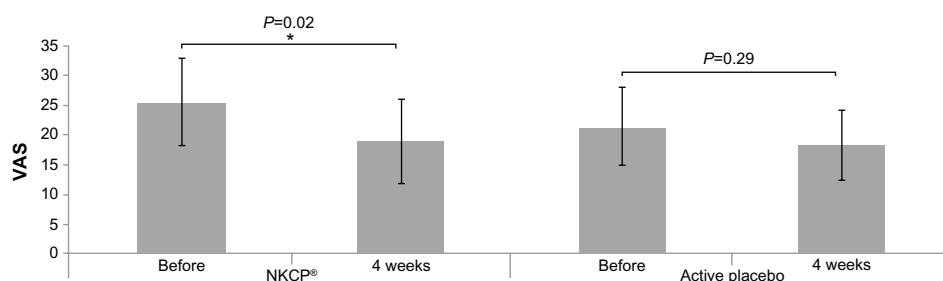


Figure 2 Changes in VAS score for low back pain after 4 weeks compared with baseline in patients taking NKCP® (Daiwa pharmaceutical Co., Ltd.) or active placebo.

Notes: The vertical bar indicates the standard error. Statistical significance, * $P<0.05$.

Abbreviation: VAS, visual analog scale.

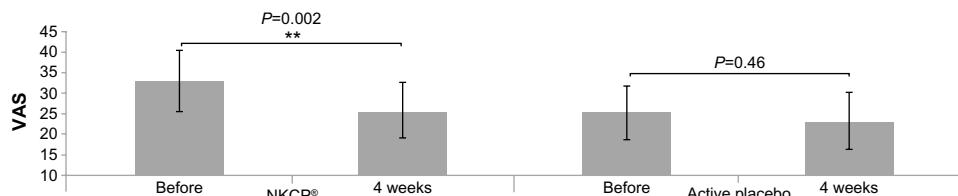


Figure 3 Changes in VAS score for coldness of the extremities after 4 weeks compared with baseline in patients taking NKCP® (Daiwa pharmaceutical Co., Ltd.) or active placebo.

Notes: The vertical bar indicates the standard error. Statistical significance, ** $P<0.01$.

Abbreviation: VAS, visual analog scale.

with increasing severity of the disease.¹⁰ In the present study, although the routine blood examination results revealed no marked abnormalities, many patients complained of low back pain, headache, shoulder stiffness, and coldness of the extremities, all of which may be partly explained by blood flow disturbances. Low back pain is an epidemiological problem that is common in the general population, and low back pain is the most dominant reason for frequent visits to general practitioners.¹¹ Although the cause of low back pain is not fully understood, atherosclerotic disease of the feeding arteries of the lumbar spine has been considered to be a possible underlying factor; additionally, a diminished blood supply is reportedly closely related to low back pain.^{12,13} In this study, improvements in the blood supply by NKCP® accordingly led to the relief of low back pain. Furthermore, the results of this study suggest that other symptoms of blood flow disturbances, such as shoulder stiffness and coldness of the extremities, also improved by the intake of NKCP®. Although statistical significances were observed in changes of these symptoms, as sample size was too small, further researches may confirm these results.

Regarding the results of blood tests, some blood chemistry parameters had abnormal values because subjects were patients of lifestyle diseases in the present study. However, the patients were well-controlled with medical intervention and had no advanced thrombotic or vascular diseases. Although a 4-week course of NKCP® significantly increased

the cholinesterase and HbA_{1c} levels, and a 4-week course of active control significantly increased the platelet count and cholinesterase level, all of these parameters were within the reference ranges, and changes were small. Therefore, we believe that these differences did not affect the results.

Various medications are used to reduce the above-described symptoms. However, because of limitations and concerns such as availability, cost, and adverse effects of medications, a proportion of the population, especially in Asia, has turned to complementary and alternative medicine.^{14,15} Japan authorities have recommended a diet balanced with staple foods, mainly Japanese foods, to maintain and improve patients' health status and to prevent chronic noncommunicable diseases.¹⁶ Therefore, functional food which prevents the development of such chronic diseases has become attractive to many people. NKCP®, which is derived from Japanese traditional food, is one of the functional foods. Direct fibrinolytic and serine protease activity for human blood and antithrombotic activity in rats after administering orally were confirmed for NKCP®.^{5,6} Furthermore, blood pressure-lowering effects for the patients have been found previously⁷ and also were shown in the present result. Therefore, NKCP® has been considered to contribute to the prevention of vascular events in patients with lifestyle diseases. In this study, we suggested that Japanese functional food improved subjective symptoms, owing to the disturbance of blood flow. As functional food is derived from Japanese

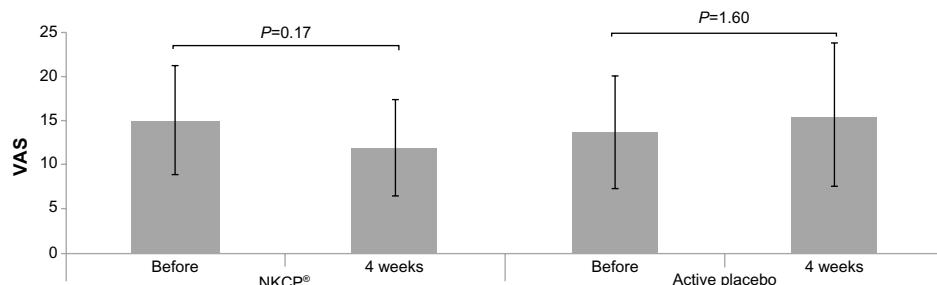


Figure 4 Changes in VAS score for headache after 4 weeks compared with baseline in patients taking NKCP® (Daiwa pharmaceutical Co., Ltd.) or active placebo.

Note: The vertical bar indicates the standard error.

Abbreviation: VAS, visual analog scale.

traditional food, natto, and does not contain other chemical substances, it is considered as safe. Thus, NKCP® may be useful for patients instead of using medications.

The most important limitation of this study is that sample size is small with a short observation period. However, as this study was performed as a cross-over, double-blind trial, there is some scientific reliability. Another limitation is that the participants of the present study were all women. The outcome of the recruitment process for participants coincidentally yielded an all-woman subject pool of participants. Further research may clarify the whether similar results can be obtained for male patients.

Conclusion

For patients with lifestyle diseases, taking NKCP® improved the blood flow and subsequently relieved low back pain, shoulder stiffness, and coldness of the extremities, all of which were caused by blood flow disturbances. The use of dietary supplements derived from the Japanese traditional food natto provides additional effects of relieving subjective symptoms for patients with lifestyle diseases who are receiving medical care. Further researches with a large enough sample size may confirm the present results.

Disclosure

This study was financially supported by Daiwa Pharmaceutical Co, Ltd. MH received a lecture fee from Daiwa Pharmaceutical Co, Ltd. KH and KM report no conflicts of interest in relation to this work. The authors report no other conflicts of interest in this work.

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