Note

Antihypertensive Effects of Nicotianamine from Soybean Broth in Spontaneously Hypertensive Rats

Tetsuo Takenaka1*, Takashi Murayama1, Tadasu Furusho2 and Yoko Takenaka3

1 Faculty of Agriculture, Tamagawa University, 6-1-1, Tamgawa-gakuen, Machida, Tokyo 194-8610, Japan
2 Department of Nutrition, Jr. College of Tokyo University of Agriculture, 1-1-1 Sakuragaoka, Setagaya-ku, Tokyo 156-8502, Japan
3 T&T Food Institute, 555-20 Aina, Astugi, Kanagawa 243-0038, Japan

Received December 1, 2008; Accepted May 11, 2009

The effect of the ACE inhibitor nicotianamine (NA), from soybean broth (SB), on blood pressure was investigated in spontaneously hypertensive rats (SHR) upon single and long-term administration. The IC50 value of NA from SB was 0.69 μmol/L. Single oral dose of NA (0.9 mg, 4.5 mg and 9.0 mg/kg body weight) decreased blood pressure 1 h after administration, and blood pressure returned to the control level 3 h after administration. Long-term oral dose of NA (0.9 mg and 4.5 mg/kg body weight) decreased blood pressure for 4 weeks after administration, while that of NA (9.0 mg/kg body weight) was decreased for the full 8-weeks feeding period. At 8 weeks after administration, serum NA content in SHR was determined by amino acid analyzer and revealed that NA was not detected in the blood of SHR (0.9 mg and 4.5 mg/kg body weight group), while 32.6 ± 7.3 μg/dL NA was detected in the 9.0 mg/kg body weight group. It was suggested that NA absorbed from the intestine decreased the systolic blood pressure (SBP) in SHR, and an appropriate NA level (9.0 mg/kg body weight group) may provide long-term antihypertensive effects upon administration.

Keywords: ACE inhibitor, antihypertensive factor, nicotianamine, soybean broth (SB), spontaneously hypertensive rats (SHR), angiotensin system

Introduction

The miso, soysauce and natto manufacturing processes produce large amounts of spent residue (soybean broth (SB)) as a waste by-product, and its disposal poses severe environmental challenges. SB contains a large amount of proteins and water-soluble polysaccharides and has a high COD (chemical oxygen demand) value, on average about 20,000 ppm, resulting in high waste-water treatment costs. The food industry’s utilization of SB as a food material is still in the developmental stage. The effective utilization of SB (Matsuda and Ueda, 1995; Kimura et al., 1997), such as the production of flavorful vinegar by fermentation (Morimura et al., 2005) and the production of GABA by lactic acid bacteria (Furuta et al., 2008), has been reported. We have already reported a method by which the ACE inhibitor nicotianamine is easily separated from SB (Takenaka et al., 2009).

The angiotensin I-converting enzyme (ACE; EC3.4.15.1) catalyzes the hydrolysis of angiotensin I to generate the potent vasoconstrictor, angiotensin II. This enzyme plays an important role in the renin-angiotensin system to regulate both arterial blood pressure, and salt and water balance. ACE inhibitors, such as captopril and enalapril, have been used as antihypertensive drugs (Gray et al., 1995).

In recent years, many ACE inhibitory peptides have been isolated from food protein sources such as casein (Maruyama et al., 1982), tuna (Kohama et al., 1988) and bonito (Matsui et al., 1993). There have been several reports about the non-peptidyl ACE inhibitors polyphenol (Horie et al., 1996) and nicotianamine (Kinoshita et al., 1993; Shimizu et al., 1999; Hayashi et al., 2005).

In this paper, we investigated the influence of nicotianamine on SBP in SHR upon single and long-term administration.
Materials and Methods

Experimental animals and measurement of blood pressure
Male SHR (11 weeks old) were purchased from Charles River Japan, and were maintained under the following conditions: temperature, 23 ± 1°C; relative humidity 55 ± 5%; 12 h light/dark cycle. All animals were allowed a standard diet (AIN-76, Oriental Yeast Industry, Tokyo, Japan) and tap water. The systolic blood pressure (SBP) was measured using a tail-cuff with a SBP monitor for rats and mice, Model NK-2000 (Muromachi Kikai Ltd., Tokyo). All the animal experiments were conducted in compliance with the guidelines of the Japanese Association for Laboratory Animal Science (1987) and guidelines for Animal Experiments of the Research and Development Division of Tamagawa University.

Materials  Soybean broth (SB), a by-product of the Natto manufacturing process, was obtained from Oyama Tofu Co. (Japan) and contained protein (8.9%), fat (3.2%), carbohydrate (77.2%), ash (10.6%) per dry weight. ACE from rabbit lung acetone powder was obtained from Sigma Chemical Co. (USA).

Hippuryl-L-histidyl-L-Leucine (Hip-His-Leu) was obtained from the Peptide Institute (Osaka, Japan). Amberlite IR-120 was obtained from Rohm & Hass Co (USA). Sephadex G-15 was obtained from GE healthcare (USA). The nicotianamine standard was kindly supplied by Dr. Kinoshita of Kikkoman Co. (Japan).

Assay of ACE inhibitory activity  ACE inhibitory activity was assayed by a modified method of Cushman and Cheung (1971). The reaction mixture was as follows: 250 μl of 12.2 mM Hip-His-Leu and 608 mM NaCl in 130 mM borate buffer (pH 8.3), 30 μl of water or inhibitor, 100 μl of ACE solution (obtained from 0.5 g rabbit lung acetone powder was obtained from Sigma Chemical Co. (USA).

Hippuryl-L-histidyl-L-Leucine (Hip-His-Leu) was obtained from the Peptide Institute (Osaka, Japan). Amberlite IR-120 was obtained from Rohm & Hass Co (USA). Sephadex G-15 was obtained from GE healthcare (USA). The nicotianamine standard was kindly supplied by Dr. Kinoshita of Kikkoman Co. (Japan).

Assay of ACE inhibitory activity  ACE inhibitory activity was assayed by a modified method of Cushman and Cheung (1971). The reaction mixture was as follows: 250 μl of 12.2 mM Hip-His-Leu and 608 mM NaCl in 130 mM borate buffer (pH 8.3), 30 μl of water or inhibitor, 100 μl of ACE solution (obtained from 0.5 g rabbit lung acetone powder was obtained from Sigma Chemical Co. (USA).

After incubation at 37°C for 1 h, the reaction was stopped by adding 250 μl of 1 M HCl. The hippuric acid released was extracted with 2.0 ml of ethyl acetate.

Quantitative analysis of nicotianamine  Concentrations of the ACE inhibitor nicotianamine in SB and serum were determined by an amino acid analyzer (JEOL JLC-500/V).

Preparation of the ACE inhibitor from SB  The ACE inhibitor was prepared by a previously described method (Takenaka et al., 2009).

Nicotianamine was isolated from SB using an ion-exchange column, Amberlite IR-120-B and Sephadex G-15 chromatography. Briefly, 300 g (dry weight) of SB was dissolved in 3 L of distilled water and centrifuged for 10 min at 1500 × g. The supernatant was applied to an Amberlite IR120-B column previously prepared as H⁺ type and then eluted with 2.7 M NH₄OH. The NH₄OH in the eluate was evaporated to dryness. The ACE inhibitor fraction from the Amberlite IR120-B column was then gel-filtered on the Sephadex G-15 column, previously equilibrated with 50 mM acetic acid, and then eluted with 50 mM acetic acid. The ACE inhibitor active fractions were pooled and lyophilized. Identification of the ACE inhibitor from SB was carried out on the IR, NMR and TOF-mass spectrometer. The ACE inhibitor active fraction was used for single and long-term administration.

Single NA administration  After preliminary breeding, 20-weeks-old male SHR were divided into four dietary groups of 6 rats each. Nicotianamine (0.9, 4.5 and 9.0 mg/kg body weight; abbreviated as follows: 0.9, 4.5 and 9.0 mg/kg groups) dissolved in saline was orally administered to SHR (3 groups), and control rats were administered an identical volume of physiological saline alone. SBP was measured every alternate hour for 5 h after ACE inhibitor administration.

Long-term NA administration  After preliminary breeding, 22-weeks-old male SHR were divided into four dietary groups of 6 rats each. As shown in Table 1, the test diets contained 0.002%, 0.010% and 0.020% of the ACE inhibitor nicotianamine were prepared by altering the proportion of or-

<table>
<thead>
<tr>
<th>Table 1. Composition of experimental diets.</th>
</tr>
</thead>
<tbody>
<tr>
<td>nicotianamine content</td>
</tr>
<tr>
<td>(mg/kg body weight)</td>
</tr>
<tr>
<td>-----------------------</td>
</tr>
<tr>
<td>Corn starch</td>
</tr>
<tr>
<td>Casein</td>
</tr>
<tr>
<td>Dextrized corn starch</td>
</tr>
<tr>
<td>Cellulose</td>
</tr>
<tr>
<td>Soybean oil</td>
</tr>
<tr>
<td>Vitamin mix (AIN76)</td>
</tr>
<tr>
<td>Mineral mix (AIN76)</td>
</tr>
<tr>
<td>L-Cystine</td>
</tr>
<tr>
<td>Choline bitartrate</td>
</tr>
<tr>
<td>sucrose nicotianamine (mg)</td>
</tr>
<tr>
<td>AIN76-MX; Oriental Yeast Co., Tokyo, Japan</td>
</tr>
<tr>
<td>AIN76-VX; Oriental Yeast Co., Tokyo, Japan</td>
</tr>
</tbody>
</table>
ordinary sucrose content. Nicotianamine was added to AIN-76 diet in amounts that corresponded to the administration of 0.9, 4.5 and 9.0 mg/kg body weight. The test diets were given to SHR for 8 weeks and after withdrawal from the test diet treatment for an additional 2 weeks.

SBP was measured weekly and body weight was measured daily. Blood was collected from SHR at 8 weeks after administration of test diets and at 2 weeks after the withdrawal from the test diets. All serum samples were kept frozen at -40°C until assayed.

Statistical analysis The statistical analysis was performed using STATCEL2 statistical software. One-factor ANOVA followed by Tukey’s test (Yanai, 2004) was used to evaluate the significance of differences among the groups. The experimental data are shown as the means ± S.E.M. Differences were considered to be statistically significant at $P < 0.05$.

Results and Discussion

ACE inhibitor concentration in SB The amount of nicotianamine in SB was 2.2 mg/g dry weight as determined by amino acid analyzer. The IC$_{50}$ value of nicotianamine from SB, as determined using rabbit lung enzyme with HHL as the substrate, was 0.21 μg/ml (0.69 μM).

Influence of nicotianamine administration on weight of rats The average body weights did not differ between groups. The average food intake and water consumption, ratios of organ weights, and SHR serum lipid levels did not differ between groups (data not shown).

Effects of single administration of nicotianamine on SBP in SHR The effects of single administration of nicotianamine on SBP in SHR are shown in Fig. 1. SBP was measured at every alternate hour after administration of test diets. The SBP was significantly lower 1 h after administration of nicotianamine (4.5 mg and 9.0 mg/kg body weight) compared to control, and returned to the control level 3 h after administration.

Kinoshita et al. (1993) reported that the HW fraction, containing nicotianamine, from soy sauce reduced SPB in SHR after oral administration. SBP was significantly decreased by 21 mmHg ($p < 0.05$) as compared to the control group 8 h after SHR were administered purified nicotianamine (100.0 mg/kg body weight), and returned to the control level at 24 h. There is a large difference in the SBP change in SHR reported by Kinoshita and our results. The dose of nicotianamine administered in our animal experiment corresponds to about 1/10 or less than that reported by Kinoshita et al. (1993). The existence of variation in the effective level of NA among these results remains to be clarified.

![Fig. 1. Effects of single administration of nicotianamine on SBP in SHR.](image-url)

Symbols: ●: control (saline at a dose of 5.0 ml/kg body weight); ■: 0.9 mg/kg group (nicotianamine at a dose of 0.9 mg/kg body weight); ▲: 4.5 mg/kg group (nicotianamine at a dose of 4.5 mg/kg body weight); ◆: 9.0 mg/kg group (nicotianamine at a dose of 9.0 mg/kg body weight). Each value is expressed as mean ± S.E.M. ($n = 6$). Significant difference from the control group: $^*P < 0.05$. 

Antihypertensive Effects of Nicotianamine from Soybean Broth in Spontaneously Hypertensive Rats 543
Effects of long-term administration of nicotianamine on SBP in SHR

The composition of test diets used for long-term administration is shown in Table 1. The SBP of the 0.9, 4.5 and 9.0 mg/kg groups were significantly lower than in the control for 2-4 weeks after NA administration initiation (Fig. 2). Subsequently, SBP levels of the 0.9 and 4.5 mg/kg groups gradually returned to the control level 6 to 8 weeks after administration initiation. However, the SBP of the 9.0 mg/kg group was significantly lower than in the control, 23.4 ± 3.4 mmHg (p < 0.05) decrease, for 8 weeks of the test feeding period. Although a sufficient antihypertensive effect was not shown at a dose of 9.0 mg/kg body weight upon single administration, an antihypertensive effect was seen upon long administration of the identical dose.

Shimizu et al. (1999) reported on the effects of long-term administration of the G fraction (containing NA) from Ashitaba and captopril on SHR. The SBP of G fraction (21.8 mg/kg body weight)- and captopril (0.3 mg/kg body weight)-treated groups were 200 ± 7.3 mm Hg (n=7) and 198 ± 8.1 mmHg (n = 5) at the end of 10 weeks of administration, respectively. The SBP of the control group was 211 ± 3.7 mmHg (n = 7). The content of NA in the G fraction remains to be clarified.

The reversal of an antihypertensive effect in the 0.9 mg and 4.5 mg/kg groups may be a result of the low ACE inhibitor affinity for vascular tissue and the high ACE inducing ability of the vascular tissue (Miyazaki, 1995). The serum nicotianamine concentration was measured using an amino acid analyzer to assess the distribution of administered nicotianamine at 8 weeks after administration and 2 weeks after withdrawal from test diet. Chromatograms of nicotianamine in serum from control diet and test diet groups at 8 weeks after administration are shown in Fig. 3. Nicotianamine was not detected in the serum of the 0.9 mg/kg and 4.5 mg/kg groups, while it was detected (32.6 ± 7.3 μg/dL; n = 6) in the serum of the 9.0 mg/kg group. Nicotianamine was not detected in the serum at 2 weeks after the withdrawal from test diet. Hayashi et al. (2005) reported that after nicotianamine was administered into the stomach of Tsukuba hypertensive mice (THM), it was absorbed from the intestine and into the blood, and consequently, the absorbed nicotianamine decreased SBP and ACE activity in the plasma. It was thought that a similar distribution of nicotianamine had been carried out in SHR, although serum ACE activity was not measured in our experiment. We are currently conducting further experiments to clarify the ACE activity and concentration of nicotianamine in the serum of SHR upon long-term administration.

![Fig. 2. Effects of long-term administration of nicotianamine on SBP in SHR. Symbols: ● control (nicotianamine at a dose of 0.0 mg/kg body weight); ■ 0.9 mg/kg group (nicotianamine at a dose of 0.9 mg/kg body weight); ▲ 4.5 mg/kg group (nicotianamine at a dose of 4.5 mg/kg body weight); ◆ 9.0 mg/kg group (nicotianamine at a dose of 9.0 mg/kg body weight). The test diet was administered for 8 weeks and its withdrawal for an additional 2 weeks. Each value is expressed as mean ± S.E.M (n = 6). Significant difference from the control group: * P < 0.05.](image-url)
Antihypertensive Effects of Nicotianamine from Soybean Broth in Spontaneously Hypertensive Rats

Fig. 3. Chromatograms of serum nicotianamine levels in control diet and test diet groups at 8 weeks after administration. Nicotianamine was measured by amino acid analyzer. Chromatogram of SHR serum in control (A) and test diet group (B) at 8 weeks after administration.

References


