Effect of Long-Term Diabetes on Serotonin-Mediated Contraction in Carotid Arteries from Streptozotocin-Induced Diabetic Male and Female Rats

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Note

An accumulating body of evidence suggests that males and females differ in vascular function in arteries under pathophysiological states. In this study, we tested whether there was a sex difference associated with serotonin (5-hydroxytryptamine, 5-HT)-mediated contraction in the carotid arteries of long-term streptozotocin (STZ)-induced diabetic rats [viz. 23 or 24 weeks after STZ (65 mg/kg, intravenously (i.v.)) injection starting at 8 weeks old of rats]. In the control group, the 5-HT- and high-K⁺-induced contractions were greater in females than in males. In both sexes, treatment with STZ led to a decrease of 5-HT-induced contraction in carotid arteries compared to controls. In STZ-induced diabetic rats, the carotid arterial 5-HT-induced contraction was greater in female rats than in diabetic male rats. The high-K⁺-induced contraction was greater in diabetic female rats than in either age-matched female controls or diabetic male rats. Expression of the 5-HT₂A receptor, which is the main receptor for 5-HT-induced contraction in rat carotid arteries, was similar among the four groups. These results suggest that decreased 5-HT-induced carotid arterial contraction is seen in both sexes under long-term STZ-induced diabetic conditions. Further, this reduction seems to be weaker in females than in males. This alteration of 5-HT-induced contraction may be partly associated with increased voltage-dependent Ca²⁺ channel activity.

Key words carotid artery; contraction; serotonin; sex difference; streptozotocin

Although women generally develop cardiovascular diseases several years later than men, this benefit may be diminished in diabetic individuals. The results of numerous epidemiological studies investigating the relationship between sex and the development of cardiovascular disease in diabetes have been inconsistent.1,2) Alterations in the responsiveness of blood vessels to various hormones and/or neurotransmitters in patients of both sexes with diabetes mellitus are well established.3–6) However, there is confounding evidence of vascular function between diabetes of both male and females.

The neurotransmitter serotonin [5-hydroxytryptamine (5-HT)] is an important factor that is involved in the regulation of several vascular functions, including blood flow, blood pressure, and vascular tone, in pathophysiological states.7–9) Several reports by our research team10–12) and others13–15) have found that alterations in vasoconstriction induced by 5-HT were observed in patients with cardiovascular diseases as well as in those with diabetes. An accumulating body of evidence suggests that 5-HT plays a role in the development of diabetic complications. For example, sarpogrelate, an antagonist of the 5-HT₂A receptor, has beneficial effects against diabetic nephropathy, neuropathy, and diabetes-associated vascular dysfunction, including arterial stiffness and arterio-sclerosis, in diabetic patients and animal models.16–19) This relevant evidence suggests that the manipulation of 5-HT function can represent a critical therapeutic target in the case of diabetic vasculopathies. In the carotid artery, which supplies blood to the brain, we found increased 5-HT-induced contractions in type 2 diabetic Goto–Kakizaki rats.20) Moreover, we very recently demonstrated that exposure to high insulin levels, but not high glucose levels, can increase 5-HT-induced contraction in rat carotid arteries.20) However, at present, whether there are sex-associated differences in carotid arterial contraction induced by 5-HT in chronic-stage diabetes remains unclear.

Type 1 diabetes can be induced by the injection of streptozotocin (STZ), which causes β-cell death in the pancreas and is commonly used to induce type 1 diabetes in experimental models.21) The aims of our study were to investigate whether 5-HT-induced carotid arterial contraction would differ between males and females with STZ-induced diabetes.

MATERIALS AND METHODS

Animals Male and female Wistar rats (8 weeks old) were randomly divided into diabetic and non-diabetic (control) groups. Experimental type 1 diabetic rats were induced with a single intravenous injection of STZ (65 mg/kg dissolved in citrate buffer), as described previously.22–24) As a control, age-matched rats were injected with citrate buffer alone. All animals were given a standard laboratory diet and water ad libitum until the rats were 31 or 32 weeks old (viz. 23 or 24 weeks after STZ/buffer injection). This study was approved by the Hoshi University Animal Care and Use Committee, and all experiments were performed in accordance with “Guide for the Care and Use of Laboratory Animals” published by the U.S. National Institutes of Health and “Guide for the Care and Use of Laboratory Animals” adopted by the Committee on the Care and Use of Laboratory Animals of Hoshi University.

Measurement of Blood Parameters and Blood Pressure

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Plasma (nonfasting; taken at sacrifice) parameters and blood pressure (measured at 1 week before sacrifice) were measured as reported previously.25–27 Blood glucose was measured by a glucose meter (OneTouch Ultra, LifeScan, Johnson & Johnson Company, Milpitas, CA, U.S.A.). Plasma lipid parameters and plasma insulin were measured using commercially available kits (Wako Pure Chemical Industries, Ltd., Osaka, Japan and Shibayagi, Gunma, Japan, respectively). Systolic blood pressure was measured by the tail-cuff technique (model BP-98A; Softron, Tokyo, Japan).

**Functional Study** Vascular isometric force of the common carotid artery was recorded as described previously.20,28,29 To generate concentration–response curves in modified Krebs–Henseleit solution ([in mM] 118.0 NaCl, 4.7 KCl, 25.0 NaHCO3, 1.8 CaCl2, 1.2 NaH2PO4, 1.2 MgSO4, and 11.0 glucose) and 5-HT (serotonin hydrochloride, Sigma-Aldrich, St. Louis, MO, U.S.A.) (10–8–3×10–3 m) or high-K+ (10–80 m vs male control, †p<0.05). In this study, long-term diabetes induced the alteration of 5-HT-induced contraction in the arterial vessels of diabetes. In this study, long-term diabetes induced the alteration of 5-HT-induced contraction in the arterial vessels of diabetes. In this study, long-term diabetes induced the alteration of 5-HT-induced contraction in the arterial vessels of diabetes. In this study, long-term diabetes induced the alteration of 5-HT-induced contraction in the arterial vessels of diabetes. In this study, long-term diabetes induced the alteration of 5-HT-induced contraction in the arterial vessels of diabetes.

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<tr>
<th>Table 1. Values of Various Parameters in STZ-Induced Diabetic and Control Male and Female Rats</th>
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<tbody>
<tr>
<td></td>
<td>Male</td>
</tr>
<tr>
<td></td>
<td>Control</td>
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<tr>
<td>Body weight (g)</td>
<td>613.2±9.4</td>
</tr>
<tr>
<td>LV/BW (mg/g)</td>
<td>1.62±0.03</td>
</tr>
<tr>
<td>SBP (mmHg)</td>
<td>105±4</td>
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<tr>
<td>HR (beats/min)</td>
<td>332.1±10.2</td>
</tr>
<tr>
<td>Glucose (mg/dL)</td>
<td>111.8±3.3</td>
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<tr>
<td>Insulin (ng/mL)</td>
<td>1.82±0.3</td>
</tr>
<tr>
<td>T-Chol (mg/dL)</td>
<td>115.8±6.5</td>
</tr>
<tr>
<td>TG (mg/dL)</td>
<td>205.6±23.1</td>
</tr>
<tr>
<td>Ring weight (mg)</td>
<td>0.93±0.05</td>
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<tr>
<td>T-Chol (mg/dL)</td>
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<tr>
<td>TG (mg/dL)</td>
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<td>Ring weight (mg)</td>
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</table>

Values are the mean±S.E. Number of experiments is shown in parentheses. LV; left ventricle, BW; body weight, SBP; systolic blood pressure, HR; heart rate, T-Chol; total cholesterol, TG; triglyceride. *p<0.05 vs. male control, †p<0.05 vs. female control, ‡p<0.05 vs. male diabetic.
diabetic female Wistar rats. Van Buren et al. \(^{32}\) observed that in mesenteric resistance arteries in short-term (4 weeks) and long-term (40 weeks) STZ (40 mg/kg, i.v.)-induced diabetic rats, the 5-HT-induced contraction was similar to age-matched control rats; however, the 5-HT-induced contraction increased and decreased in the basilar artery in short-term and long-term STZ-induced diabetic rats, respectively. These discrepancies may result from vessel types, sex, or duration of disease.

The STZ-induced diabetic method is commonly used to generate type 1 diabetic animal models. \(^{21,33}\) In this study, male rats treated with a single dose of STZ exhibited lower BW, hyperglycemia, and hypoinsulinemia than controls; however, the 5-HT-induced contraction increased and decreased in the basilar artery in short-term and long-term STZ-induced diabetic rats, respectively. These discrepancies may result from vessel types, sex, or duration of disease.

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In the present study, we also found increased levels of blood lipid parameters (namely, total cholesterol and triglyceride) in the diabetic group compared with those in the control group in each sex, and the alteration of reactivity to 5-HT was seen in hyperlipidemic conditions. \(^{39,40}\) Because the above blood parameters did not differ between male and female diabetic rats, these factors may not be associated with 5-HT-induced contraction in carotid arteries. Moreover, we very recently found that high insulin exposure, but not high glucose levels, led to enhanced 5-HT-induced contraction in rat carotid arteries. \(^{20}\) Therefore, we suggest that the insulin level is a determinant factor in contractile response to 5-HT in carotid arteries. Further investigation is required to identify causative factors in the alteration of 5-HT-induced vasocontraction between sexes.

Vascular contractile mechanisms are mainly divided into two mechanisms: calcium signals and changes in the contractile apparatus’ sensitivity to calcium. \(^{41,42}\) Indeed, these two signaling pathways are utilized in 5-HT\(_{2A}\) receptor-mediated responses in vascular smooth muscle. \(^{9,11–14,20,43}\) In the present study, high-K\(^+\)-induced contraction in diabetic female rats was greater than in control female rats and diabetic male rats. We suggest that the greater response to 5-HT in diabetic female carotid arteries relative to that of diabetic males may be partly attributable to increased voltage-gated calcium channel activity rather than 5-HT\(_{2A}\) receptor expression. Because insulin...
can modulate intracellular calcium levels, the different insulin levels under diabetic conditions between sexes may also contribute to altered high-K+-induced contraction. However, further investigation is required to establish the mechanisms underlying these factors. In conclusion, we suggest that decreased 5-HT-induced carotid arterial contraction is observed in both sexes under long-term STZ-induced diabetes and the extent of the reduction differs between sexes. Our findings provide evidence of sex differences related to reactivity to 5-HT in long-term diabetic states and may contribute to the development of new strategies for the treatment of diabetes-associated vascular dysfunction.

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Conflict of Interest The authors declare no conflict of interest.

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