TRYPTAMINERGIC MECHANISM PARTICIPATING IN INDUCTION OF VASOCONSTRICTION BY ADENINE NUCLEOTIDES, ADENOSINE, IMP AND INOSINE IN THE ISOLATED AND BLOOD-PERFUSED HINDLIMB PREPARATION OF THE RAT

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Abstract—The isolated right hindlimb of the recipient rat was perfused at a constant flow rate through the femoral artery with heparinized blood from the carotid artery of a donor. The preparations were under a 99.0 ± 0.8 mmHg of mean perfusion pressure (N = 63) and 3.3 ± 0.1 ml/min of blood flow through the right femoral artery. The actions of adenosine, adenosine tri-, di- and monophosphate, inosine monophosphate and inosine on the femoral vascular bed were investigated, respectively. These substances injected into the femoral artery, with the exception of inosine, caused a dose-dependent vasoconstriction always preceded by a temporal vasodilatation. Inosine induced only a prompt vasoconstriction. The vasoconstrictor responses to these substances were diminished or reverted to vasodilator ones after repeated administrations and such were significantly prevented by pretreatment with either reserpine or methysergide. These results indicate that all the purines tested induce a vasoconstriction in the femoral vascular bed of the rat through a common (tryptaminergic) mechanism and that such seem to be potent releasers of 5-hydroxytryptamine from peripheral tryptaminergic storage sites.

Since Drury and Szent-Györgyi (1) first investigated the actions of adenosine and adenine nucleotides on the cardiovascular system, considerable interest has been focused on the mode of actions of these substances in the peripheral circulation. It has been reported that intra-arterial injection of these substances causes vasodilatation on the femoral vascular bed of the cat (2) and dog (3–6).

Recently, however, it was reported that either adenosine or inosine given into the femoral artery of the rat caused a prominent vasoconstriction which was associated with a tryptaminergic mechanism (7–8). Adenine nucleotides are hydrolysed in blood and tissues and converted rapidly to adenosine and/or inosine (9–11). It was of interest, therefore, to examine whether adenine nucleotides and inosine monophosphate also induce a constriction of the femoral vasculature of the rat through a tryptaminergic mechanism.

The present study, using isolated and cross-circulated hindlimb preparation of the rat, provides pharmacological evidence that adenine and hypoxanthine series release 5-hydroxytryptamine from the peripheral stores, resulting in a vasoconstriction of the femoral artery.

MATERIALS AND METHODS

Male Sprague-Dawley rats were allowed free access to food and water overnight prior
to experiments.

Surgical procedures

Donor rats (550-750 g) were anaesthetized initially with pentobarbital sodium (65 mg/kg i.p.), and anaesthesia was maintained hourly with sustaining doses of urethane (400 mg/kg s.c.). The right jugular vein and carotid artery were cannulated with polyethylene cannulae. Heparin sodium (1000 U/kg) was injected into the femoral vein, and the systemic blood pressure was measured from the left femoral artery with a pressure transducer (Nihon Kohden, RP-5). The carotid and jugular cannulae of the donor were connected to the perfusion circuit after the circuit had been filled with about 10 ml of blood freshly drawn from heparinized rats.

Recipient rats (300-350 g), which were anaesthetized with pentobarbital sodium (65 mg/kg i.p.), were prepared for perfusion of the right hindlimb. The right femoral artery and vein were dissected free, and loops of thread for ligatures were placed around the blood vessels. The muscles such as M. rectus femoris, M. psoas longus and M. gracilis were carefully ligated and cut around the caput femoris. The femoral and sciatic nerves were also severed. Then, an abdominal midline incision was made and the intestine was moved toward the left for exposure of the right common iliac artery and vein near the origin of the abdominal aorta and inferior vena cava. After heparin sodium (1000 U/kg) was injected into the left femoral vein, a venous cannula (ID 1 mm, OD 1.5 mm) was inserted into the femoral vein via the common iliac vein. Subsequently, an arterial metal cannula (ID 0.4 mm, OD 0.6 mm) was introduced into the femoral artery via the common iliac artery. Immediately after the right hindlimb of the recipient was isolated completely from the body by cutting the bone at the caput femoris, the blood from the right carotid artery of the donor was conducted into the femoral artery of the isolated right hindlimb of the recipient by means of a peristaltic pump (Mitsumi Science, SJ-1210). The pump was precalibrated and rechecked at the end of the experiment. A square wave electromagnetic flowmeter (Nihon Kohden, MF-25) was used for the measurement of the femoral blood inflow. Perfusion pressure was measured with a pressure transducer (Nihon Kohden, RP-5) from a side arm near the arterial cannula. Recordings were made on an ink-writing rectigraph (TOA Electronics, EPR-3T). The experimental set-up is illustrated schematically in Fig. 1A.

The perfusing blood from the carotid cannula of the donor first flowed into a small glass bottle. The volume of blood in the bottle was maintained constant by adjustment with a small screw clamp. The venous outflow from the femoral vein of the isolated limb of the recipient was collected in a venous reservoir and in turn returned to the jugular vein of the donor by 15 cm drop of hydrostatic pressure. The donor and the isolated right hindlimb of the recipient, respectively, were placed on heating tables and the body temperature was maintained between 36 and 38 °C throughout the experiment. Moreover, the surface of the hindlimb was covered with liquid paraffin and warmed by radiant heat from a lamp.

Reserpine was given s.c. twice in dose of 5 mg/kg 48 and 24 h, respectively, prior to the experiment.
FIG. 1. Schematic diagram of the preparation (A in the Figure). Blood flows from the carotid artery of a donor into an arterial reservoir (AR) and is constantly pumped to the femoral artery (FA) of the isolated right hindlimb of the recipient (B in the Figure). Femoral venous blood is returned via a venous reservoir (VR) to the jugular vein of the donor. BP, systemic blood pressure; Temp, thermometer; W, circulating warming water; AR, arterial reservoir; F, filter (Sartorius-Membranfilter GMBH, West Germany); Pump, peristaltic pump; BS, blood sampling; FM, electromagnetic flowmeter; PP, perfusion pressure; FA, femoral artery; FV, femoral vein.

**Drugs**

The drugs used were adenosine-5'-triphosphate disodium (ATP), adenosine-5'-diphosphate sodium (ADP), adenosine-5'-monophosphate sodium (AMP), inosine-5'-monophosphate sodium (IMP), adenosine, inosine and 5-hydroxytryptamine creatinine sulphate (5-HT) (Sigma), (±)-noradrenaline hydrochloride (Sankyo), methysergide tartrate (Sandoz), tyramine hydrochloride (Tokyo Kasei) and reserpine (Apoplon®, Daiichi Seiyaku). Drugs were dissolved in 0.9% saline as stock solutions and were diluted with 0.9% saline just before use. Microsyringes (Jintan Terumo Co.) were used for close-arterially injecting a volume of 10 μl in a period of 4 sec into a rubber tube connected to the shank of the polyethylene cannula. Doses of 5-HT, tyramine, noradrenaline and methysergide refer to their salts and of the other substances to their bases. Increases (vasoconstriction) or decreases (vasodilatation) in the perfusion pressure caused by drugs were taken as drug responses, as the perfusion rate was constant.

**Statistical analysis**

Values in the text are means±S.E. The statistical significance of the differences between mean values was analysed using Student's t-test and expressed as P values.
RESULTS

Blood flow through the femoral artery

Within 30 min after the beginning of perfusion, the mean perfusion pressure reached a constant level of 99.0 ± 0.8 mmHg (N = 63). The preparations then had a femoral blood flow of 3.3 ± 0.1 ml/min and such remained stable for 3 hr or more. The average wet weight of the perfused hindlimb measured at the end of the experiment was 29.6 ± 0.3 g, and the flow rate per unit of wet weight was 10.1 ± 0.4 ml min⁻¹ 100 g⁻¹.

Effects of ATP, ADP, AMP, IMP, adenosine and inosine on the femoral vascular bed

Single injections of increasing doses (3 × 10⁻⁹ - 10⁻⁶ moles) of either ATP, ADP, AMP, IMP or adenosine into the femoral artery produced a prominent vasoconstriction preceded...
by a temporal vasodilatation (Fig. 2A). Inosine \((3 \times 10^{-9} - 10^{-5} \text{ moles})\) induced only a vasoconstriction. The vasoconstrictor responses to these substances were increased in a dose-dependent manner in doses of \(3 \times 10^{-9} \text{ to } 3 \times 10^{-7} \text{ moles}\), and reached the maximum at \(3 \times 10^{-7} \text{ moles}\), and at \(10^{-6} \text{ moles}\) the vasoconstrictor responses tended to decrease (Fig. 2B).

Responses of the femoral vascular bed to repeated administrations of either ATP, ADP, AMP, IMP, adenosine or inosine

All the purines tested were injected into the femoral artery in an equimolar dose \((10^{-6})\). Further explanation is described in the equal for Fig. 2.

Fig. 3. Response of the femoral vascular bed to repeated administrations \((10^{-6} \text{ moles})\) of either ATP, ADP, AMP, IMP, adenosine (Ads) or inosine (Ino). Further explanation is described in the equal for Fig. 2.
mole). The first administration (I) of these substances caused a marked and long-lasting vasoconstriction. When the responses to the preceding administrations had worn off completely, the same dose (10⁻⁶ moles) of these substances was successively injected, respectively. As shown in Fig. 3A and B, the vasoconstrictor responses to these substances were diminished progressively after repeated administrations, and finally almost disappeared or vasodilator responses were seen.

When the intensity of vasoconstrictor responses to the first administrations of these substances was compared, there was statistically no significant difference (Fig. 3B).

*Vasoconstrictor potencies of ATP, ADP, AMP, IMP, adenosine and inosine in the femoral vascular bed*

As shown in Fig. 2B, all the purines tested produced dose-dependent vasoconstrictor responses within the dose range of 3 × 10⁻⁹ to 3 × 10⁻⁷ moles, but at 10⁻⁶ moles the dose-

<table>
<thead>
<tr>
<th>Derivative</th>
<th>Negative log mol ED50 (moles)</th>
<th>No. of experiments</th>
</tr>
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<tbody>
<tr>
<td>ATP</td>
<td>7.21±0.08</td>
<td>5</td>
</tr>
<tr>
<td>ADP</td>
<td>7.29±0.03</td>
<td>5</td>
</tr>
<tr>
<td>AMP</td>
<td>7.12±0.07</td>
<td>5</td>
</tr>
<tr>
<td>IMP</td>
<td>7.20±0.14</td>
<td>5</td>
</tr>
<tr>
<td>Adenosine</td>
<td>7.10±0.07</td>
<td>7</td>
</tr>
<tr>
<td>Inosine</td>
<td>7.05±0.10</td>
<td>5</td>
</tr>
</tbody>
</table>

ED50 values were obtained from the dose-response curves reconstructed in such a way that the responses to 3 × 10⁻⁷ moles of each were taken as a 100% maximum vasoconstrictor ones, and are expressed as the negative logarithm of the molar dose of the purines tested producing a 50% maximum vasoconstriction. Values are means ± S.E.

**Fig. 4.** Responses of the femoral vascular bed to tyramine (Tyr) and ATP or IMP in reserpinized preparations. A) Original tracings. Compare with the responses to the same dose of ATP or IMP in untreated preparations (see Fig. 3A). B) Summarized data. Open columns: untreated; hatched columns: treated with reserpine. The results of IMP (N=2) are not shown. N.S., not significant. Other details as in Fig. 2.
response curves fell. Thus, it was reasonable to consider that these substances consistently caused a 100% maximum constriction of the femoral vasculature at the dose of $3 \times 10^{-7}$ moles.

The ED50 (the negative logarithm of the molar dose producing a 50% maximum vasoconstriction) was determined by reconstructing dose-vasoconstrictor response curves for each substance in a percentage of the maximum response (Table 1). There was no significant difference in the potencies of these substances inducing the vasoconstrictor responses.

**Effect of pretreatment with reserpine on the vasoconstrictor responses to ATP and IMP**

A single injection of ATP (500 $\mu$g; $10^{-6}$ moles) or IMP (350 $\mu$g; $10^{-6}$ moles) was given to a single preparation. Although the vasoconstrictor response to the first single injection (I) of either ATP or IMP into the femoral artery in reserpine-pretreated animals was not significantly different from the vasoconstrictor response to the same dose of ATP or IMP in untreated preparations (Fig. 3A and B), the vasoconstrictor response to the second injection (II) was diminished to a greater extent than that in untreated preparations (Fig. 4A and B). The response to tyramine (30 $\mu$g) was considerably sustained in reserpinized
preparations.

**Abolition by methysergide of the vasoconstrictor response to either ATP or IMP**

As high doses of ATP or IMP which produce overt responses induced a tachyphylaxis, only a single dose of each was administered to a single preparation. In this connection, in untreated preparations (Fig. 5A) noradrenaline (0.1 μg), 5-HT (1 μg) and ATP (500 μg; 10⁻⁶ moles) or IMP (350 μg; 10⁻⁶ moles) were given in that order and responses to these substances served as controls. The effect of methysergide was examined in other preparations in which noradrenaline and 5-HT were given prior to methysergide, and noradrenaline, 5-HT and ATP or IMP were injected after methysergide in that order. As shown in Fig. 5A, a single injection of methysergide (1 μg) into the femoral artery, which completely inhibited the vasoconstrictor response to 5-HT (1 μg) but not that to noradrenaline (0.1 μg), had almost no effect on the perfusion pressure. With the same dose of methysergide, the vasoconstrictor response to either ATP or IMP was almost completely blocked (Fig. 5A and B).

**DISCUSSION**

The present study revealed that when injected into the femoral artery of the rat, adenine nucleotides and IMP as well as adenosine and inosine caused prominent vasoconstrictions and that these responses showed marked tachyphylaxis. In the same sort of preparations, it has already been suggested that adenosine and inosine induce a vasoconstrictor response probably by releasing 5-HT from the peripheral stores, on the basis of the facts that the response disappeared in both of the reserpine- and methysergide-treated preparations (7-8).

ATP is rapidly decomposed to adenosine or inosine via AMP or IMP by dephosphorylation in blood and tissues (9-11). It would be reasonable to suggest, therefore, that all the purines tested act as 5-HT releasers inducing vasoconstriction in the hindlimb of the rat. In fact, the vasoconstrictor responses to ATP and IMP, which were picked up out of six purines tested, were less prominent in reserpinized preparations as compared with that to the same doses of these substances in untreated preparations. Furthermore, the treatment with methysergide definitely blocked the constriction of femoral vasculature induced by ATP or IMP as well as 5-HT without preventing the vasoconstrictor effect of noradrenaline. These findings indicate that ATP and IMP as well as adenosine and inosine are potent releasers of 5-HT, regardless of the fact that the site from which 5-HT is released remains to be determined.

It should be emphasized that in order to release 5-HT from the peripheral storage site these substances must primarily enter the cell. However, a high phosphate compound such as ATP, unlike adenosine and inosine, does not readily cross cell membranes (12-13). It seems likely, therefore, that exogenous nucleotides are broken down to adenosine or inosine before taken up, in view of the evidence that nucleosides can readily pass through cell membranes (12, 14).

The present results did not show a statistically significant difference between the potencies of these substances inducing vasoconstriction. This can be explained on the basis that
nucleotides are rapidly degraded to adenosine or inosine in blood and tissues.

From this study and the previous reports (7-8), it was concluded that all the purines tested produced a definite vasoconstriction indirectly by releasing 5-HT from the peripheral stores, and that the intrinsic vasodilator responses to these substances were probably masked by the predominant vasoconstrictor ones.

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