Effects of serotonin on the internal anal sphincter: *in vivo* manometric study in rats

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SUMMARY The effects of serotonin (5-hydroxytryptamine, 5-HT) on the internal anal sphincter were studied in anaesthetised rats. Serotonin induced a dose dependent relaxation of the internal anal sphincter. Methysergide blocked this relaxation, but did not affect the rectoanal reflex. Methysergide did not antagonise the actions of cholinergic and adrenergic agonists on the internal anal sphincter. Other 5-HT antagonists such as cyproheptadine, ketanserin, chlorpromazine, amitriptyline and ergotamine failed to reduce or block the internal anal sphincter relaxation due to 5-HT, nor did they alter the rectoanal reflex. Adrenergic and cholinergic antagonists had no effect on the 5-HT-induced relaxation of the internal anal sphincter, suggesting that 5-HT acts upon the internal anal sphincter *via* a non-adrenergic, non-cholinergic mechanism. Tetrodotoxin reduced or blocked the relaxation of the internal anal sphincter produced by 5-HT, implying that 5-HT acts through neural pathways rather than directly on the muscle. It is concluded that although 5-HT relaxes the internal anal sphincter, it does not participate in the rectoanal reflex.

Serotonin (5-HT) has been proposed as a neurotransmitter in various parts of the alimentary tract acting on both enteric nerves and on the smooth muscles.¹ Pharmacological studies of 5-HT have revealed its multiple actions in the gut. Responses of the smooth intestinal muscle to 5-HT are quite variable, depending on the region of the gut and the species examined.^{2,3} Serotonin acts on enteric ganglia,^{4,5} thereby activating neurones that release acetylcholine and cause contraction of the longitudinal and circular layers of smooth muscle.^{6,7}

Serotonin also activates non-adrenergic, non-cholinergic intrinsic inhibitory neurones, and is therefore able to relax the gut if excitatory neuromuscular transmission is blocked.⁶⁻⁸ These results suggest that the role of 5-HT in the enteric nervous system is to serve as the transmitter of interneurones.¹ Its effect on the gut motor activity have been both excitatory and inhibitory, probably by stimulating different receptor systems.^{9,10}

Burleigh *et al*¹¹ reported that 5-HT induced contraction of the human internal anal sphincter *in vitro*, and therefore excluded it as a possible inhibitory neurotransmitter. Rayner,¹² however,

reported that 5-HT relaxed the internal anal sphincter muscle strip of the vervet monkey in organ bath.

Serotonin has been implicated in disturbances of gastrointestinal motility in patients with carcinoid syndrome. In a study of patients with Hirschsprung's disease, Rogawski *et al*¹³ reported the absence of serotonergic neurones in the aganglionic colon.

The pharmacology of the internal anal sphincter has been studied *in vivo* using animal models.^{12,14} No information is available, however, on the actions of 5-HT on the internal anal sphincter *in vivo*. The aim of this study has been to investigate the effects of 5-HT and its antagonists on the tone of the anus and rectum and the rectoanal reflex in intact rats.

Methods

RATS

One hundred and five rats of either sex were anaesthetised with ip chloral hydrate 200 mg/kg. The carotid artery and jugular vein were cannulated using a 19G needle catheter. Blood pressure was measured with a transducer connected to the carotid artery and drugs were administered through the jugular vein. Anal and rectal pressures were recorded with a probe containing a proximal rectal balloon and a distal anal balloon which was posi-

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tioned within the lumen of the internal anal sphincter.¹⁵ The rectoanal reflex was elicited by inflating the rectal balloon with 0.6–0.8 ml air. Pressures were recorded *via* Statham pressure transducers by a Hewlett-Packard 7745A recorder. When needed, a respirator was used to maintain adequate ventilation. The administration of each drug was repeated several (five to 10) times in a given experiment. There was sufficient time between injections to allow for full recovery.

Drugs used were: serotonin (creatinine sulphate complex), tryptamine hydrochloride, cyproheptadine hydrochloride, amitriptyline hydrochloride, 1,1-dimethyl 4-phenylpiperazinium iodide (DMPP), carbamylcholine chloride (carbachol), isoprenaline hydrochloride, noradrenaline bitartrate, hexamethonium bromide and tetrodotoxin (TTX), all from Sigma. Methysergide dimaleate (Sandoz), phenoxybenzamine hydrochloride (SK&F), propranolol hydrochloride (Abic), atropine sulphate (Teva), ketanserin (Janssen).

Results

EFFECTS OF 5-HT AND TRYPTAMINE

Injection of serotonin (5-HT) in the range of 3–50 $\mu\text{g/kg}$ relaxed the internal anal sphincter in all 105 rats (Fig. 1). The magnitude of the relaxation is expressed by multiplying the response amplitude by its duration and is dose-dependent (Fig. 2). In the

Fig. 1 Actions of serotonin (5-HT) on the rectum (RECT) and internal anal sphincter (IAS); pressure recording in the rat. Inflation of the rectal balloon with 0.7 ml of air, produced IAS relaxation – the recto-anal reflex. 5-HT relaxed the IAS and contracted the rectum in a dose dependent manner. Note the decrease in blood pressure (BP) caused by 5-HT. Concentrations are given in $\mu\text{g/kg}$.

Fig. 2 Dose dependent curves for the 5-HT-induced relaxation of the IAS, obtained in 13 rats. The 5-HT effect is expressed by multiplying the response amplitude ($\text{cm H}_2\text{O}$) by its duration (min) and it is dose dependent. Note that in spite of the absolute variations, the threshold and range of the responses are similar for most of the experiments.

rectum, 5-HT caused contraction (70% of rats) which was likewise dose dependent (Fig. 1). Repeated 5-HT doses of up to 100 $\mu\text{g/kg}$ (accumulated dose), did not change the amplitude or duration of the relaxation of the internal anal sphincter, compared with a control response.

Tryptamine (25–1000 $\mu\text{g/kg}$) elicited a dose dependent relaxation of the internal anal sphincter (five rats) and was 50–100 times less potent than 5-HT in producing internal anal sphincter relaxation. This relation was determined by injecting 5-HT and tryptamine at varying doses and then comparing the doses needed to produce similar responses (Fig. 3). The blood pressure was transiently reduced after 5-HT administration. In contrast, tryptamine caused an increase in blood pressure followed by a decrease. Similar observations have been made in several animal species, and it has been suggested that 5-HT and tryptamine act *via* different receptors.¹⁶

EFFECTS OF 5-HT ANTAGONISTS

Methysergide (32 $\mu\text{g/kg}$), reduced the effect of 5-HT upon the internal anal sphincter by about 50% (Fig. 4). At higher doses (200 $\mu\text{g/kg}$, $n=12$) it completely blocked the 5-HT-induced relaxation of the internal anal sphincter (Fig. 4). It did not alter the resting

Fig. 3 Comparison of the effect of tryptamine (TRYP) and 5-HT on the IAS. Increased doses of tryptamine, up to 500 µg/kg, produced IAS relaxation and increase in blood pressure (BP). 5-HT was 50–100 times more potent than tryptamine in producing IAS relaxation. The effects of both 5-HT and tryptamine on the IAS, were blocked by methysergide.

tone of the rectum and anus and the rectoanal reflex (Fig. 4).

The effects of cholinergic and adrenergic agonists such as: DMPP, carbachol, phenylephrine and isoprenaline upon the internal anal sphincter were not affected by methysergide (200–400 µg/kg). Administration of other 5-HT antagonists: cyproheptadine (up to 4 mg/kg, $n=7$), ketanserin (up to 6 mg/kg, $n=6$), chlorpromazine (up to 0.1 mg/kg, $n=4$), amitriptyline (up to 0.6 mg/kg, $n=3$) and ergotamine (up to 0.25 mg/kg, $n=4$) failed to affect the relaxation of the internal anal sphincter induced

by 5-HT. The anal and rectal tones and the rectoanal reflex were also not affected.

EFFECTS OF ADRENERGIC AND CHOLINERGIC ANTAGONISTS

The actions of 5-HT may be mediated by the release of other neurotransmitters such as noradrenaline or acetylcholine. To test this possibility, adrenergic and cholinergic antagonists were given before 5-HT administration. The alpha adrenoceptor antagonist phenoxybenzamine (250 µg/kg) blocked the effect of noradrenaline (1.5 µg/kg, $n=7$) but did not affect the internal anal sphincter relaxation induced by 5-HT (Fig. 5a).

The beta adrenoceptor antagonist propranolol (1 mg/kg, $n=5$) blocked the effect of isoprenaline (6 µg/kg) on the internal anal sphincter, but failed to affect the relaxation of the internal anal sphincter produced by 5-HT (Fig. 5b). The muscarinic cholinergic antagonist atropine (0.5 mg/kg, $n=3$) blocked the effect of carbachol (2 µg/kg) but did not change the relaxation of the internal anal sphincter to 5-HT administration (Fig. 6). Hexamethonium, a nicotinic cholinergic antagonist, (2.4 mg/kg, $n=5$) blocked the effect of DMPP (60 µg/kg) upon the internal anal sphincter, but had no effect on the relaxation of the internal anal sphincter produced by 5-HT (Fig. 6b).

EFFECT OF TETRODOTOXIN

Receptors for 5-HT have been found on both muscle

Fig. 4 Methysergide antagonism of 5-HT actions on the IAS. Methysergide (METHYS) at a low dose of 32 µg/kg reduced the 5-HT-induced IAS relaxation. At a dose of 200 µg/kg methysergide completely eliminated the response to 5-HT. During this blockade of 5-HT effects, the rectoanal reflex remained unchanged. Phenylephrine (PE) was given to show that the IAS muscle retains its ability to relax in the presence of methysergide.

Fig. 5 Effects of adrenergic antagonists on the IAS response to 5-HT: (a) the effect of alpha adrenergic blockade. Noradrenaline (NA) caused brief relaxation of the IAS. Phenoxybenzamine (PHENOXY) antagonised the effect of NA but did not alter the response to 5-HT. (b) Effect of beta adrenergic blockade. Isoprenaline (ISOP) caused a prolonged relaxation of the IAS, which was blocked by propranolol (PROPRAN). The response to 5-HT was not affected by propranolol.

and nerve cells. To distinguish between these two sites of action, the nerve blocker tetrodotoxin was used. Tetrodotoxin, in an accumulated dose of 2.8 to 18.4 μg/kg was applied to eight rats. In all eight rats tetrodotoxin blocked or reduced the internal anal sphincter relaxation to 5-HT administration in a dose dependent pattern (Fig. 7). As shown in Fig. 7, a dose of 3 μg/kg of tetrodotoxin completely blocked the effect of 5-HT (3 and 6 μg/kg) on the internal anal sphincter, while it reduced the internal anal sphincter relaxation induced by 9 μg/kg of 5-HT. In order to block the effect of higher 5-HT doses (10–40 μg/kg) increased doses of tetrodotoxin (10–18.4 μg/kg) were used.

Tetrodotoxin blocked the internal anal sphincter response to DMPP (60 μg/kg) which is nerve mediated, while it had no effect upon internal anal

Fig. 6 Effect of cholinergic antagonists on the IAS response to 5-HT: (a) muscarinic blockade by atropine (ATROP). Carbachol (CARB) contracted the IAS and the rectum. After the application of atropine this contraction was blocked, but the 5-HT response was not affected. (b) nicotinic blockade by hexamethonium (HEXAMETH). The nicotinic cholinergic agent DMPP induced IAS relaxation. Hexamethonium blocked this effect but did not change the response to 5-HT.

sphincter relaxation produced by phenylephrine (14 μg/kg) which presumably acts directly on the smooth muscle (Fig. 7).

Discussion

Serotonin (5-HT) displays a multiplicity of actions as a potential neurotransmitter in the gastrointestinal tract.^{1 17 18} In the present *in vivo* study in rats, the internal anal sphincter underwent relaxation by 5-HT. This finding correlates with the *in vitro* study by Rayner¹² who found that the internal anal sphincter of the vervet monkey is relaxed by 5-HT. In contrast to the present results, Burleigh *et al*¹¹ reported that 5-HT induced contraction of the human internal anal sphincter in organ bath. The reason for this difference is not clear; it may be

Fig. 7 Actions of tetrodotoxin (TTX) on the IAS relaxation evoked by 5-HT. Effects of DMPP, which are nerve mediated, were blocked by TTX whereas phenylephrine (PE) which acts directly on the muscle was still effective. During nerve blockade by TTX the relaxations caused by 5-HT (3 and 6 $\mu\text{g/kg}$) were completely prevented and the response to 5-HT (9 $\mu\text{g/kg}$) was greatly reduced. Note the increase in the spontaneous activity of the IAS after TTX injection.

because of the species difference, or because of the experimental conditions (*in vitro* vs *in vivo*). It should be mentioned that Burleigh *et al* did not specify the site of 5-HT actions in the human internal anal sphincter, which may be the muscle itself. Our experiments on the rat indicate that 5-HT acts *via* a neural pathway.

In the present work, 5-HT produced in the majority of cases (70%), a dose dependent rectal contraction. Our results show that 5-HT was 50–100 times more potent than tryptamine in producing internal anal sphincter relaxation. Methysergide was the only effective antagonist, while ketanserin and cyproheptadine failed to block 5-HT actions. This indicates that the receptors mediating 5-HT actions in the internal anal sphincter may be similar to those described as 5-HT₁ central receptors.¹⁹ The relaxation of the internal anal sphincter after 5-HT administration was completely blocked by methysergide, which failed to affect the rectoanal reflex. It may therefore be concluded that 5-HT does not act as an inhibitory neurotransmitter in the rectoanal reflex.

It has been proposed that methysergide at high concentrations is comparably effective in antagonising the actions of carbachol, DMPP and histamine on the intestine.¹⁷ The effect of methysergide on cholinergic and adrenergic agonists activity was examined, using the same concentrations that blocked the internal anal sphincter response to 5-HT. Methysergide had no effect upon the internal anal sphincter response to administration of: carbachol,

DMPP, isoprenaline and phenylephrine and seems therefore to act as a selective 5-HT antagonist.

It has been shown that during exposure of isolated intestinal segments to 5-HT, the sensitivity to this drug is diminished.⁷ The effect is dependent on the presence of 5-HT in the organ bath and it is reversed on washing. Such a tachyphylactic effect has not been observed in our experiments. A possible explanation for this finding is that in the intact animal, the 5-HT level is readily reduced by the effective uptake mechanism for this substance.^{1, 17}

Serotonin can stimulate⁷ or antagonise²⁰ the release of acetylcholine in the gut. This raises the possibility that 5-HT acts on the internal anal sphincter by modifying the release of acetylcholine or perhaps of other transmitters such as noradrenaline. Adrenergic and cholinergic agents were shown to modify the activity of the rat internal anal sphincter.²¹ To examine this point, cholinergic and adrenergic antagonists were used. Administration of such antagonists (atropine, hexamethonium, phenoxybenzamine and propranolol) failed to affect the internal anal sphincter relaxation produced by 5-HT, as well as the tone of the anus and rectum. This shows that the action of 5-HT is not mediated by cholinergic or adrenergic transmitters.

Methysergide may antagonise muscular^{6, 10} as well as neural^{20, 22} 5-HT receptors. In order to eliminate the possible neural effect of 5-HT, tetrodotoxin was used. Tetrodotoxin blocked or reduced the effect of 5-HT on the internal anal sphincter. This finding suggests that 5-HT causes internal anal sphincter

relaxation *via* activation of neural rather than muscular receptors. A similar effect of tetrodotoxin on 5-HT neural receptors in the gut was observed by other investigators.^{7,12,20,23} It also seems reasonable to conclude that methysergide blocks the 5-HT effect by acting at the neural level. The latter conclusion is supported by the studies of Wood and Mayer²² and North *et al.*²⁰ who observed the direct effect of methysergide on myenteric neurones.

In conclusion, the present results indicate that 5-HT stimulates a non-adrenergic, non-cholinergic inhibitory neural pathway with a consequent relaxation of the internal anal sphincter in rats. It is not the neurotransmitter, however, that causes the relaxation of the internal anal sphincter during the rectoanal reflex.

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