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A comparison of the hormone treated groups however, shows that progesterone does not reduce the tension in response to electrical stimulation. Similarly, there is no significant difference between the oestrogen- and the progesterone-dominated uteri in the degree of calcium binding. These results contrast with similar studies in the rabbit (Schofield, 1955; Csapo, 1956) and pig (Knifton, 1966).

The only effect of progesterone on the rat myometrium that this study reveals therefore, is a decrease in the sensitivity to oxytocin.

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References

Bengtsson, L. Ph. & Schofield, B. M. (1960). J. Reprod. Fert., 1, 402-409.
Csapo, A. (1950). Am. J. Physiol., 162, 406-410.
Csapo, A. (1956). Recent Progr. Horm. Res., 12, 405-427.
Csapo, A. & Corner, G. W. (1952). Endocrinology, 51, 378-385.
Knifton, A. (1966). J. Pharm. Pharmac., 18, Suppl., 151S-159S.
Schofield, B. M. (1955). J. Physiol., Lond., 129, 289-304.
Schofield, B. M. (1960). Ibid., 154, 48P-49P.
Schofield, B. M. (1963). In Recent Advances in Physiology, editor Cresse, R., pp. 222-251, London: Churchill.

Nature of adrenergic receptors on the skin melanophores of Rana tigrina

SIR,—This report describes the experiments conducted to determine the nature of adrenergic receptors on frog melanophores.

Adult *Rana tigrina* (80–350 g) were anaesthetized by injection of pentobarbitone sodium (50 mg/kg) into the abdominal cavity. Drugs, dissolved in 0.6% saline or amphibian Ringer solution, were injected through the cannulated left branch of thoracic aorta (Bhide & Gupta, 1967) or through one of the liver lobes. Skin colour was observed with the naked eye, and melanophores of the web skin were graded by the method of Hogben & Slome (1931). In experiments in conscious frogs, drugs dissolved in distilled water were injected into the abdominal cavity. From 3 to 6 frogs were used for each dose of each drug and the average change in melanophore index recorded (Tables 1 and 2).

Noradrenaline, which acts predominently on α -type adrenergic receptors was more potent in concentrating melanin in melanophores than adrenaline which

TABLE 1	. EFFECT	OF	ADRENERGIC	DRUGS	ON	THE	MELANOPHORE	INDEX J	IN	Rana
	tigrina									

	Anaesthe	etized frogs	Conscious frogs			
Drug	Dose mg/kg (No. of frogs)	Average change in the melanophore index Decrease (-) Increase (+)	Dose mg/kg (No. of (frogs)	Average change in the melanophore index Decrease (-) Increase (+)		
Noradrenaline hydrochloride .	. 0·1 (7) 0·5 (7) 1·0 (5)	-1.2 -1.8 -2.6	1·0 (4) 4·0 (9)	-2·0 -3·5		
Adrenaline hydrochloride	. 0.5 (3) 1.0 (7)	-2.6 -0.75 -1.25 -1.8	0·3 (5) 1·0 (3) 3·0 (3)	-0.7 -1.33 -2.33		
Isoprenaline sulphate	. 2.0 (5) 0.3 (5) 1.0 (3) 3.0 (4)	$ \begin{array}{c} -1.8 \\ +0.67 \\ +1.0 \\ +0.88 \end{array} $	3.0 (3) 0.3 (3) 1.0 (4) 3.0 (6)	$ \begin{array}{c} -2.33 \\ +0.66 \\ +1.0 \\ +1.5 \end{array} $		

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TABLE 2. EFFECT OF ADRENERGIC BLOCKING DRUGS ON NORADRENALINE-INDUCED CHANGE IN MELANOPHORE INDEX IN *Rana tigrina*. Phenoxybenzamine was given 40 min and other blocking drugs 5–10 min before noradrenaline

	Anaesthe	tized frogs	Conscious frogs			
Blocking agent (dose mg/kg)		Dose of noradrenaline mg/kg (No. of frogs)	Average decrease in melanophore index	Dose of noradrenaline mg/kg (No. of frogs)	Average decrease in melanophore index	
None (taken from Table 1) Phenoxybenzamine (25.0)		$ \begin{array}{r} 1 \cdot 0 (5) \\ 1 \cdot 0 (3) \\ 2 \cdot 0 (4) \\ 6 0 (4) \end{array} $	2.6 0.0 0.2 1.25	1.0 (4) 2.0 (4) 6.0 (3)	2·0 0·0 0·0	
Dihydroergotamine methane- sulphonate (1.0) Yohimbine hydrochloride (1.5)	•••	6·0 (6) 1·0 (6) 6·0 (4) 1·0 (7)	0.5 1.6 0.6	0.0 (3)	0.0	
Phentolamine methanesulphonate (0.25) . Pronethalol hydrochloride* (2.0) INPEA hydrochloride* (2.0) .	 	1 · 0 (4) 1 · 0 (4) 1 · 0 (4)	0·38 1·0 1·62	1·0 (4) 1·0 (5)	1.9 1.6	

* These drugs themselves caused blanching and decreased melanophore index by 0.5 to 2.0. Further decrease induced by noradrenaline is given here.

acts on both α - and β -receptors (Table 1). Phenoxybenzamine, dihydroergotamine, phentolamine and yohimbine, which block α -adrenergic receptors, blocked or much reduced the action of an effective dose (1 mg/kg) of noradrenaline (Table 2). This confirms previous reports that ergotamine (Lerner, 1959) and phenoxybenzamine (Bhide & Gupta, 1967) block the action of adrenaline and noradrenaline on frog melanophores.

On the other hand, isoprenaline produced dispersion of melanin and this action was blocked by β -adrenergic blocking agents. Pronethalol and INPEA (*N*-isopropyl-*p*-nitrophenylethanolamine), themselves caused some concentration of melanin and did not block further action of noradrenaline on melanophores.

The present work suggests that the adrenergic receptors on skin melanophores of *Rana tigrina* are predominently of the α -type and they are responsible for colour change induced by adrenaline and noradrenaline. It also suggests occurrence of β -type adrenergic receptors on the melanophores.

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References

Bhide, N. K. & Gupta, I. (1967). J. Pharm. Pharmac., 19, 58-59. Hogben, L. & Slome, D. (1931). Proc. R. Soc., Lond., 108B, 10-53. Lerner, A. B. (1959). Nature, Lond., 184, 674-677.