Indian J Physiol Pharmacol 1995; 39(1): 86-88

ROLE OF 5-HYDROXYTRYPTAMINE IN TOXAEMIA OF PREGNANCY

T. K. BHATTACHARYYA* AND P. K. DEBNATH**

*Department of Pharmacology, School of Tropical Medicine, Calcutta - 700 073

and

**Department of Internal Medicine, J.B. Roy State Ayurvedic Medical College Hospital, Culcutta - 700 004

(Received on September 15, 1993)

Abstract: 5-HT concentration in blood of 24 randomly selected pregnant women presenting with signs and symptoms of toxacmia of pregnancy were estimated and compared with 30 normal pregnant women. As such 5-HT level increases with the progress of pregnancy and in toxacmia group in comparison to normal pregnancy. A change in the degree of oedema from moderate to severe could bring a statistically significant change in blood pressure and serum 5-HT level. Increased 5-HT plays a role in toxacmia as well as in normal pregnancy and some of the toxic effects observed in toxacmia of pregnancy are due to the effects of 5-HT.

Key words : toxaemia of pregnancy

5-HT

serotonin

INTRODUCTION

The presence of serotonin (5-Hydroxytryptamine, 5-HT) in the mammalian reproductive system (1) had stimulated investigators to find out its role in the normal physiological functions and in diseased state in obstetrics. Attempts have been made to implicate 5-HT in toxaemia of pregnancy (2) also, although Israel (3) failed to observe such role. In certain studies relating to 5-HIAA (a metabolic product of 5-HT) excretion (3) in urine of normal vis-a-vis toxaemia of pregnancy, no definite correlation with 5-HT and its excretion in normal as well as toxaemia of pregnancy (3, 4) could be established. A conclusive role of 5-HT in maintainance and termination of pregnancy could not be established as none of available data relating uterine and placental 5-HT content in human being (1) could show the values at different stages of pregnancy in any systematic manner.

This prompted the authors to study the role of 5-HT in normal and toxaemia of pregnancy.

METHODS

The study was conducted on 30 normal pregnant mothers and 24 patients with different grades of toxacmia attending antenatal O.P.Ds at J.B. Roy State Ayurvedic Medical College Hospital, Calcutta and S.S. K.M. Hospital, Calcutta. The age ranged from 18 to 24 years and the subjects were otherwise healthy with average built and nutrition and of different socioeconomic status. The criteria to grade the toxacmic Indian J Physiol Pharmacol 1995; 39(1)

REFERENCES

- Bikhazi GB, Leung I, Foldes FF. Interaction of neuromuscular blocking agents with calcium channel blockers. *Anaesthesiology* 1982; 57; A 268.
- Bikhazi GB, Leung I, Foldes FF. Calcium channel blockers increase potency of neuromuscular blocking agents in vivo. Anaesthesiology 1983; 59: A276.
- Chattopadhyay RN, Roy RK, Das AK. Comparative studies of the effects of calcium channel blockers on isolated skeletal muscle preparation. *Indian J Pharmacol* 1992; 24 : 233-234.
- Godfriend T, Miller R, Wibo M. Calcium antagonism and calcium entry blockade. *Pharmacol Rev.* 1986; 94/1: 321-416.
- Kondo K, Suzuki H, Okuno T et al. Effects of nifedipine, diltiazem and verapamil on vasoconstrictor responses to norepinephrine and potassium ion in rat mesenteric artery. Arch Int Pharmacodyn 1980; 245 : 211-219.
- 6. Schwartz A, Triggle DJ. Cellular action of calcium antagonist

drugs. Annual Review Med 1984; 35: 325-389.

- Wang T, Tsali LI, Schwartz A. Effects of calcium antagonists on sarcoplasmic reticulum. *Europ J Pharmacol* 1984; 100: 253-261.
- Burn JH. In : practical pharmacol, Blackwell Scientific Publication, Oxford 1952.
- Ghosh MN. In fundamentals of Exp. Pharmacol, Calcutta Scientific Agency 1971: 56-57.
- Zclis R, Flaims SF. Calcium influx blockers and vascular smooth muscle; do we really understand the mechanism? *Annals Intern Med* 1981; 94 : 124-136.
- 11. Banergy, Levis. Physiol Pharmacol 1969, vol. III, p. 104.
- Thorens S, Hausler G. Effects of some calcium channel inhibitors on calcium translocation in vascular smooth muscle. *Europ J Pharmacol* 1979; 54 : 79-91.

Indian J Physiol Pharmacol 1995; 39(1)

patients were based on blood pressure and oedema. The patients were grouped according to the period of gestation : 32-34 wks and 35-39 wks. The period of gestation was calculated from the first day of the last menstrual period. Blood samples were analysed for 5-HT spectrofluorometrically following the method of Snyder et al (5).

RESULTS

The blood 5-HT concentration (μ g/ml) in normal and toxaemia of pregnancy in different

5-HT in Toxaemia of Pregnancy 87

DISCUSSION

Reports suggesting increase in 5-HT concentration with the progress of pregnancy (6) and more so with toxaemia of pregnancy (2) and the demonstrated ability of increased 5-HT to induce foetal death and deformity in pregnant rats and mice with carcinoid syndrome (4) hints at a strong possible role of 5-HT in the pathophysiology of gestation including toxaemia of pregnancy. Also, an increase in 5-HT level in toxaemia (4, 7) and antagonizing effect of specific 5-HT inhibitors (2) in pregnancy, suggest a possible

TABLE 1: 5-HT concentration $(\mu g/m)$ in normal and toxagenia of pregnancy in diffe	nt periods of gestation.	
---	--------------------------	--

mg/ml/	Blood 5-IIT concentration (Mean ± S. E. M.)			
Veek of gestation	<i>(n)</i>	normal	<i>(n)</i>	Toxaemia
32-34	15	0.289±0.032	12	0.398±0.059
35-39	15	0-331±0-041	12	0.439±0.052
Pooled	30	0·297±0·038	24	0-423±0-065

TABLE II : Relationship between blood 5-HT concentration and blood pressure with grades of oedema in toxaemia of pregnancy. (Mean \pm S. E. M.)

Degree of oedema	(n)	Blood pressure (mm of Hg)		Serum 5-HT (µg/ml)
		Systolic	Diastolic	
Control	24	120.63 ± 3.95	78.55 ± 3.49	0.308 ± 0.027
Moderate (++)	14	138.67 ± 4.97*	$101.33 \pm 4.65*$	0.365 ± 0.036
Severe (+++)	10	159.33 ± 5.02**	$128.53 \pm 5.64**$	0.509 ± 0.042**

P value v/s control : * < 0.01, ** < 0.001

gestational periods (32 - 34 and 35 - 39 wks) are not significantly different although an apparent increase was observed in toxaemia groups (Table I). The relationship between blood 5-HT concentration and blood pressure with oedema is shown in Table II. Worsening of oedema seems to play a significant role in blood pressure as well as blood 5-HT content. influence of 5-HT in toxaemia of pregnancy. 5-HT regulates certain physiological functions and some of its effects in toxaemia and normal pregnancy are direct. Imbalance in the production of vasoactive PGs (thromboxane A_2 and prostacyclin) (8) leading to the vasoconstriction of small arteries and platelet activation observed in toxaemia of pregnancy, may also be 5-HT mediated, atleast to a significant extent like some other physiological functions widely discussed.

Bhattacharyya and Debnath

88

Indian J Physiol Pharmacol 1995; 39(1)

REFERENCES

- Koren Z, Pfeifer Y, Sulman FG. Serotonin content of human placenta and fetus during pregnancy. Am J Obstet Gynec 1965, 93:411-415.
- Senior JB, Fahim I, Sullivan FM, Robson JM. Possible role of 5-hydroxytryptamine in toxaemia of pregnancy. *Lancet* 1963; 2:553-554.
- Israel SL, Seligson HT, Stroup PE, Seligson D. Serotonin in pregnant and parturient women. *Obstet Gynecol* 1959; 13: 672-676.
- Robson FM, Sullivan FM. Effect of 5-hydroxtryptamine on maintainance of pregnancy, congenital abnormalities and the development of toxaemia. Adv Pharmacol 1968; 6B: 187-189.

- Snyder SH, Axelrod J, Zweig M. Sensitive and specific fluorescence assay for tissue serotonin. *Biochem Pharmacol* 1965; 14:831-835.
- Koren Z, Pfeifer Y, Sulman FG. Induction of legal abortion by intrauterine instillation of Pargyline hydrochloride (Eutonyl). J Reprod Fert 1966; 12: 75-79.
- Carter FB, Cherny WB, GreenShaw C. Serotonin studies in abnormal pregnancies. Am J Obstet Gynec 1962; 84 : 913-918.
- Walsh SW. Pre-eclampsia : an imbalance in placental prostacyclin and thromboxane production. Am J Obstet Gynec 1985; 152 : 335-340.

1. No secondary, in classes of THE Consequential because participation in classes of the Consequence southers participation formula to all southers and rO's artitized or the aminophilicity of a southership of a subscript for the participation of the southership of a subscript of the participation of the southership of a subscript of the participation of the southership of a subscript of the participation of the southership of a subscript of the participation of the southership of a subscript of the subscript of the southership of a subscript of the subscript pretational particle 11. 11 and 35. 39 what are not supplied and different children in a second difference with observed in transmiss groups (Table 1). The participation of pretation difference in shown in a difference with orderent corner to shown in a considered pretation are well as blood or shifted in blood pretation as well as blood a difference.