

# Endometrial oxytocin binding sites in normal women and in subfertile patients

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**Summary:** Specific binding of oxytocin to high affinity sites in endometrial membrane preparations has previously been shown in sheep. Endometrial tissue preparations from 27 'normal' women of proven fertility were incubated with tritiated oxytocin and the existence of significant binding sites in human endometrium was shown. Furthermore, the level of binding sites underwent a cyclical variation with the highest concentration of binding at midcycle. A cyclical pattern of binding site concentration not unlike that found in the normal women was shown in 19 subfertile patients taking clomiphene.

However, in 20 subfertile patients not taking clomiphene, no cyclical pattern emerged with significantly lower levels of binding sites in the mid-portion of the cycle ( $P < 0.02$ ) and significantly higher levels in the mid-late luteal phase ( $P < 0.01$ ), as compared to the normal women. In the mid-portion of the cycle levels were significantly lower in the subfertile patients not taking clomiphene ( $P < 0.03$ ) as compared to those taking clomiphene. No significant differences were shown between the normal women and those patients taking clomiphene.

## Introduction

Although one or more causes can now be identified for many cases of subfertility, there remains a substantial proportion of patients whose primary subfertility is unexplained.<sup>1</sup> It is possible that some of these women may have subtle defects in the endometrium that do not enable the tissue to respond adequately to hormones secreted by normally functioning ovaries. Indeed, a defect in the ability of cells to take up progesterone into the nucleus<sup>2</sup> and a significantly lower level of cytoplasmic progesterone receptors have been described in endometrium from such women.<sup>3</sup> Progesterone receptor synthesis is a product of normal oestradiol activity, and it has been shown in rabbit uterine tissue that synthesis of specific binding sites for oxytocin is also stimulated by oestradiol, and repressed by progesterone.<sup>4</sup>

Oxytocin is produced and secreted by the posterior pituitary gland and this is involved, through high affinity binding sites in the mammary glands<sup>5</sup> and myometrium,<sup>6</sup> in the processes of lactation and parturition. Recently, however, it has been shown that the corpus luteum, at least in the ewe, is capable of synthesizing oxytocin<sup>7</sup> and thus may play a role in the normal control of fertility by action on oxytocin receptors which have been found to be present in the endometrium.<sup>8</sup> Oxytocin

has been shown to stimulate the production of luteolytic PGF<sub>2</sub><sup>9</sup> and during luteal regression in sheep, intermittent surges of PGF<sub>2</sub> secretion from endometrium are accompanied by similar surges of plasma oxytocin.<sup>10</sup> In view of the close relationship of oxytocin receptor concentrations to the steroid hormones, we have studied the binding of oxytocin to endometrial material in both fertile and subfertile women in an attempt to understand more about the role of the peptide hormone in the human.

## Patients and methods

### Patients

Endometrial tissue was obtained by dilatation and curettage from 66 women attending the University Hospital, Nottingham. Normal tissue was obtained from 27 patients admitted for sterilization who had had at least one full-term pregnancy. They were not taking steroidal contraceptives nor using an intrauterine contraceptive device. Tissue was also obtained from 39 patients with primary subfertility. All 39 patients had a normal pelvis at laparoscopy with fallopian tube patency as demonstrated by bilateral spill of dye. Consort's semen analysis was within the laboratory's quoted normal limits. Nineteen of the 39 patients were taking the anti-oestrogen drug clomiphene. In all 66 patients serum oestradiol and progesterone levels were

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consistent with stated day of cycle, as was histological dating of endometrial tissue.

Part of the curettage material was put in formal saline for histological examination, and the remainder frozen at  $-20^{\circ}\text{C}$ . Peripheral blood was taken from every patient within 24 hours of operation and serum oestradiol and progesterone concentration measured.

### Methods

All procedures were performed at  $4^{\circ}\text{C}$ .

**Oxytocin uptake** Endometrial samples were rinsed in ice-cold 0.9% (w/v) NaCl containing EDTA (1 mmol/l). The material was then homogenized in 25 mmol/l tris-HCl buffer (pH 7.6) containing 0.25 mol/l sucrose and 1 mmol/l EDTA. Homogenates were centrifuged at  $200 \times g$  for 10 minutes, the resulting supernatants being further centrifuged for 1 hour at  $95000 \times g$ . The pellets obtained were washed with tris-HCl buffer and then re-suspended by homogenization in 500 ml of the same buffer.  $^3\text{H}$ -Oxytocin binding in the subcellular fractions thus produced was measured by incubating 100  $\mu\text{l}$  of the fraction at  $25^{\circ}\text{C}$  for 15 minutes in 100 ml tris-HCl buffer containing 0.1% (w/v) bovine serum albumin, 2 mmol/l  $\text{MnCl}_2$  and 60 nmol/l  $^3\text{H}$ -oxytocin. Non-specific binding was measured in the presence of 200  $\mu\text{mol}$  unlabelled oxytocin. After incubation, 2 ml ice-cold tris-HCl buffer containing 1% (w/v) bovine serum albumin was immediately added. The particulate fraction was then removed on to fine glass fibre filters and washed twice with 2 ml tris-HCl buffer containing 0.1 (w/v) bovine serum albumin. The filters, containing membrane-bound  $^3\text{H}$ -oxytocin, were then counted in a liquid scintillation counter. The specific binding was calculated from the difference in binding (per mg protein) between the incubation with and without excess oxytocin. The protein in 100 ml of each endometrial sample was measured by the method of Lowry *et al.*<sup>11</sup> using bovine serum albumins as standards.

**Serum steroids** Serum oestradiol and progesterone were measured by specific radioimmunoassays with inter-assay coefficients of variation of 8.8% and 6.5% respectively at median concentrations. The normal ranges in the laboratory are: progesterone in the follicular phase 0.32–1.9 nmol/l and luteal phase 6.4–7.9 nmol, oestradiol in the follicular phase 73.4–257 pmol/l, at midcycle 367–1468 pmol/l and in the luteal phase 184–1101 pmol/l.

**Statistical analysis** In calculations of statistical significance Fisher's exact probability test was used for binding site concentrations and the Student *t*-test for the serum steroid concentrations.

### Results

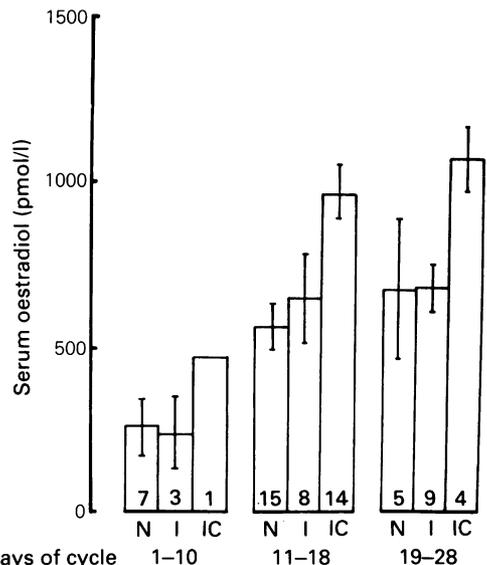
The mean ages of the three groups  $\pm$  s.e.m. were normal women  $33.4 \pm 0.8$  years, subfertile women not taking clomiphene  $28.6 \pm 0.6$  years, subfertile women taking clomiphene  $28.2 \pm 0.6$  years.

Figures 1 and 2 indicate the serum steroid concentration (mean values  $\pm$  s.e.m.) in normal women (N), subfertile patients not taking clomiphene (I) and subfertile patients taking clomiphene (IC). The number of patients in each group is indicated at the base of the respective columns.

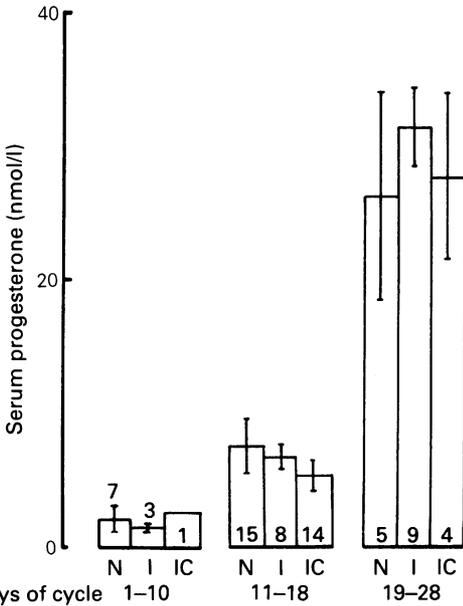
The serum oestradiol concentrations of patients in the mid-portion of their menstrual cycle (days 11–18) were found to be significantly higher in patients on clomiphene as compared to normal women ( $P < 0.001$ ) and to subfertile patients not taking clomiphene ( $P < 0.001$ ). Serum oestradiol concentrations in patients on clomiphene were also significantly higher in the mid/late secretory phase of the cycle (days 19–28), as compared to normal women ( $P < 0.01$ ) and to subfertile patients not taking clomiphene ( $P < 0.001$ ). No differences were found in progesterone levels.

Figures 3–5 illustrate the levels of oxytocin binding site concentration plotted against the day of menstrual cycle, defined as time from the beginning of last menstrual period.

When the concentration of oxytocin binding in 'normal' women was compared to that in patients with unexplained primary subfertility not taking



**Figure 1** Mean serum oestradiol concentrations  $\pm$  s.e.m. in normal women (N), subfertile patients not taking clomiphene (I) and subfertile patients taking clomiphene (IC). Figures at base of column indicate number of patients in each group.



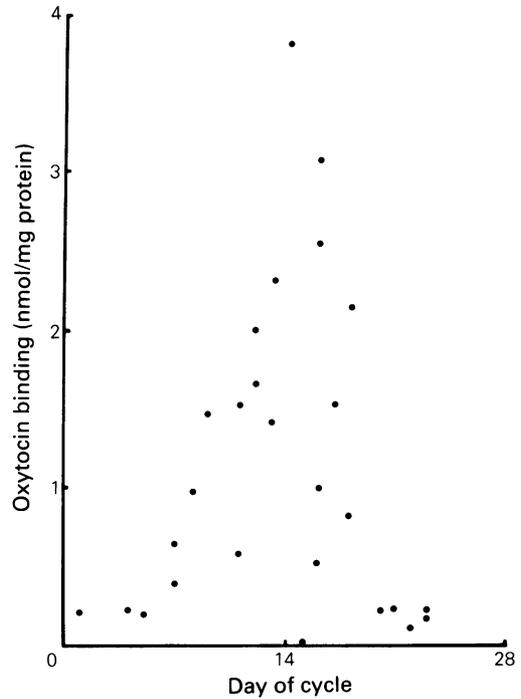
**Figure 2** Mean serum progesterone concentration  $\pm$  s.e.m. (see Figure 1 for details).

clomiphene, significantly higher levels of binding were found in the mid-portion of the cycle (days 11–18),  $P < 0.01$ . Similarly, oxytocin binding in the mid-portion of the cycle (days 11–18) in subfertile patients not taking clomiphene was significantly higher. In the mid/late luteal phase of the cycle significantly higher levels of oxytocin binding were found in subfertile patients not taking clomiphene as compared with the normal women ( $P < 0.01$ ). No significant differences were found between the normal women and the subfertile women taking clomiphene.

**Discussion**

The results obtained show that in normal women, the concentration of oxytocin binding sites displays a cyclical variation during the menstrual cycle (Figure 3). The concentration of binding sites was found to be low in the early proliferative phase (around 0.2 nmol/mg protein) and increased throughout the proliferative phase, reaching a maximum value of almost 4 nmol/mg protein at midcycle. The level then decreased during the secretory phase, reaching a value of approximately 0.2 nmol/mg protein during the mid-late secretory phase of the cycle. These findings are similar to those found in sheep endometrium by Roberts *et al.*<sup>8</sup> and Sheldrick and Flint.<sup>12</sup>

Oestradiol and progesterone endometrial binding sites have also been shown to demonstrate a cyclical variation in normal menstruating women,<sup>13</sup>



**Figure 3** Oxytocin binding sites in endometrial tissue from normal women.

with the high level of oestradiol and progesterone receptors in the late proliferative phase of the cycle correlating with raised levels of oestrogens. It is believed that oestrogens stimulate protein synthesis leading to accumulation of steroid receptors,<sup>14</sup> and that a raised level of progesterone receptors at mid-cycle is important for implantation.<sup>15</sup> Depressed levels of both steroid receptors during the secretory phase of the cycle coincide with uterine domination by progesterone which is believed to induce the down regulation of both steroid receptors.<sup>16</sup>

When the group of unexplained primary subfertile women was studied, two distinct patterns of oxytocin binding emerged. It was found that the concentration of binding sites in the subgroup of patients not taking clomiphene did not follow any recognizable pattern (Figure 4) although their plasma steroid concentration differed little from those in the normal group. However, when binding site concentrations in the subgroup of patients taking clomiphene were examined, a cyclical pattern not unlike that in the normal group was found (Figure 5).

Clomiphene therapy to subfertile women is derived from significantly increased pregnancy rates in women so treated.<sup>17</sup> Fleming and Coutts<sup>18</sup> found that clomiphene increased follicular development as indicated by elevated pre-ovulatory plasma oestrogen levels, but that this was not



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