The effects of oestradiol benzoate, progesterone, relaxin and ovariectomy on cervical extensibility in the late pregnant rat

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Summary. Cervical extensibility increased from Day 16 to term in the pregnant rat. Following ovariectomy on Day 16 of pregnancy the cervix became as inextensible by Day 20 as that of non-pregnant animals. Fetal growth was maintained in rats ovariectomized on Day 16 if given oestradiol benzoate plus progesterone but cervical extensibility only increased to a small extent. Relaxin given to these animals further increased cervical extensibility, suggesting a role for this hormone.

Introduction

Normal parturition depends upon co-ordinated uterine activity and cervical dilatation. The properties of the rat cervix show two distinct changes during pregnancy. There are increases in both the inner circumference of the cervix (associated with increased tissue mass) and its extensibility or ability to extend continuously under prolonged loading (Harkness & Harkness, 1959; Harkness & Nightingale, 1962; Rundgren, 1974). In the rat both properties increase progressively from about Day 12 until the end of pregnancy. Measurements of tensile properties of the cervix by conical probes, as used in many earlier studies, probably reflect changes in inner circumference (Harkness, 1964).

Uyldert & De Vaal (1947) showed that the inner circumference of the cervix still increased during pregnancy in rats whose uterine horns were surgically separated from the cervix, suggesting hormonal control of this change. Oestrogens alone and in combination with progesterone and/or relaxin can increase cervical size and produce small increases in cervical extensibility in non-pregnant and mid-pregnant, ovariectomized rats (Kroc, Steinetz & Beach, 1959; Cullen & Harkness, 1960; Zarrow & Yochim, 1961). Cervical dilatation in the pre-term sheep following fetal dexamethasone infusions was said to be inhibited by progesterone injections (Liggins, 1973) although this has not been supported by more quantitative measurements (Stys, Clewell & Meschia, 1978).

There is therefore evidence that the ability of the cervix to extend does increase during pregnancy but the hormonal control of this increase is less certain. The objective of the present experiments was to determine the effects of various hormonal manipulations on cervical extensibility in the late pregnant rat. Preliminary results have been published (Hollingsworth & Isherwood, 1977).

Materials and Methods

Primiparous Sprague-Dawley rats (300–450 g), of known gestational age (day of finding copulation plug = Day 1), were housed in a daily 13-h light period (07:00–20:00 h). Most of the rat in this colony and photoperiod give birth between 12:00 h on Day 22 and 12:00 h on Day 23.
After treatment of the rats (see below) they were killed, their cervices removed and cervical wet weight determined. The cervix was defined as the less vascular tissue with parallel lumina between the uterine horns and the vagina (Harkness & Harkness, 1959; Hollingsworth & Isherwood, 1978). Weight of total products of conception (fetuses, placentas, amniotic fluid and membranes), and numbers of alive and dead fetuses per litter were routinely determined.

Tensile properties of the isolated cervix were measured by a modification (Hollingsworth & Isherwood, 1977) of the method of Harkness & Harkness (1959) and Harkness & Nightingale (1962). Briefly, the cervix was mounted in a tissue bath containing Krebs–Henseleit (1932) solution at 37°C. A 60 g load was applied to a pin placed through one cervical canal while another pin, placed through the other canal, was fixed. This load produced an extension of the cervix. Readings of extension were taken every min for 5 min and then every 5 min, generally for 100 min. From these readings cervical inner circumference could be calculated (Harkness & Harkness, 1959) and a typical result is shown in Text-fig. 1.

Text-fig. 1. Effect of a 60-g load on the inner circumference of the cervix of a Day-22 pregnant rat; (a) plot of experimental values (△) and calculated line from fit of a hyperbola plus a straight line, and (b) separated into linear and hyperbolic components $I = $ initial circumference (mm) at 0 min before adding load. $l_i$ = inner circumference (mm) at 0 min obtained by extrapolation of the linear component. $K = $ rate of increase in inner circumference (mm/min) for the linear component. $T_m = $ time to half maximum for hyperbolic component. Cervical extensibility $= 10^3 \frac{K}{l_i}$ (min$^{-1}$), which is the fractional increase in circumference per min.

Inner circumference increased linearly with time after an initial curve following application of the load as described by Harkness & Harkness (1961). The mean circumference of the head of a rat fetus at Day 22 of pregnancy, determined with cotton thread on 84 fetuses, was 43.6 mm. This is approximately the circumference to which the cervical lumina must be stretched at parturition. The use of a NOVA 820 computer (Data General Corporation), a curve-fitting programme (Steven, Podraský & Foster, 1978) and data from 10 of our experiments showed that the initial curves were well fitted by a hyperbolic and less well by an exponential component. The curve-fitting programme was given starting values for four parameters, $I_h$ (circumference in mm at zero time before adding load), $I_i$ (inner circumference in mm at zero time, obtained by extrapolation of the linear component), $K$ (rate of increase in inner circumference in mm/min for the linear component) and $T_m$ (time in min to half maximum for the hyperbolic component), which will describe the algebraic sum of a hyperbola plus a straight line. The parameters were independently changed so that both hyperbolic and linear components were fitted simultaneously until the deviations of the computed points from the experimental points reached a minimum. Over 95% of experiments could be fitted within 1 mm of the experimental points (Text-fig. 1).

A derived measure of the extensibility of the cervix was used (Harkness & Harkness, 1961; Harkness & Nightingale, 1962), namely $10^3 \frac{K}{l_i}$, with units of min$^{-1}$, which is the fractional increase in circumference per min, and is independent of between-experiment variations in the size of the inner circumference ($I_i$). The measures of the hyperbolic component, $T_m$ and ($l_i$ —
were greater for the cervices of the pregnant rats than for those of the non-pregnant and post-partum rats but were not accurately measured by the current technique. The curve fit was used primarily to obtain a more accurate measurement of extensibility and inner circumference (Iₐ).

Oestradiol benzoate (3-benzoxyloxyoestra-1,3,5(10)-triene-3,17β-diol) and progesterone (pregn-4-ene-3,20-dione) were obtained from Sigma Chemical Co. and injected in arachis oil. Gifts of relaxin, NIH-R-P1 (442 i.u./mg) from NIAMDD, Bethesda, U.S.A., and CMa1-18-28AAE (3000 i.u./mg) from Dr B. G. Steinetz, were injected in saline (9 g NaCl/l).

Results are presented as means ± s.e.m. and statistical comparisons were made using the Mann–Whitney U-test (Siegel, 1956).

Results

Preliminary experiments showed that the mean ± s.e.m. extensibility (7·3 ± 1·3 min⁻¹) and inner circumference (18·2 ± 1·0 mm) of cervix of Day-18 pregnant rats after incubation for 2 h in Krebs–Henseleit solution (1932) were not different from the values of extensibility (5·8 ± 0·7 min⁻¹) and inner circumference (15·9 ± 1·1 mm) following 30 min incubation. All tissues were studied within 2 h of removal from the rats.

Experiment 1: tensile properties of the cervix during pregnancy

The cervix of the non-pregnant rat was only slightly extended during 100 min under load but extensibility was increased 8-fold by Day 22 of pregnancy (Table 1). Cervical wet weight and inner circumference increased during pregnancy with a similar time course but proportionately less. By Day 1 post partum extensibility was similar to the non-pregnant value but there was a smaller percentage decrease in wet weight and inner circumference.

Table 1. Properties of the cervix (extensibility (10³ K/Iₐ) and inner circumference (Iₐ)) and products of conception of the rat during pregnancy and after birth

<table>
<thead>
<tr>
<th>Reproductive status</th>
<th>No. of rats</th>
<th>Wet wt. (g)</th>
<th>Extensibility (min⁻¹)</th>
<th>Inner circumference (mm)</th>
<th>Weight (g)</th>
<th>Mean no. of fetuses alive/total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Non-pregnant</td>
<td>7</td>
<td>0·09 ± 0·01</td>
<td>1·3 ± 0·7</td>
<td>9·0 ± 0·5</td>
<td>12·8 ± 0·5</td>
<td>13·4/13·4</td>
</tr>
<tr>
<td>Pregnant (day)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>16·7 ± 1·1</td>
<td>13·0/13·0</td>
</tr>
<tr>
<td>16</td>
<td>8</td>
<td>0·14 ± 0·01</td>
<td>4·3 ± 1·2</td>
<td>12·4 ± 1·4</td>
<td>12·8 ± 0·5</td>
<td>13·4/13·4</td>
</tr>
<tr>
<td>17</td>
<td>4</td>
<td>0·17 ± 0·01</td>
<td>5·2 ± 1·9</td>
<td>13·3 ± 1·3</td>
<td>23·4 ± 1·1</td>
<td>12·8/13·0</td>
</tr>
<tr>
<td>18</td>
<td>8</td>
<td>0·17 ± 0·01</td>
<td>5·6 ± 1·2</td>
<td>16·2 ± 0·7</td>
<td>36·7 ± 2·6</td>
<td>12·7/12·7</td>
</tr>
<tr>
<td>19</td>
<td>7</td>
<td>0·19 ± 0·01</td>
<td>9·1 ± 1·8</td>
<td>19·7 ± 0·7</td>
<td>43·2 ± 3·7</td>
<td>11·5/11·5</td>
</tr>
<tr>
<td>20</td>
<td>6</td>
<td>0·22 ± 0·02</td>
<td>7·0 ± 0·8</td>
<td>22·2 ± 0·8</td>
<td>59·1 ± 2·7</td>
<td>11·9/11·9</td>
</tr>
<tr>
<td>21</td>
<td>7</td>
<td>0·23 ± 0·01</td>
<td>9·1 ± 1·3</td>
<td>22·9 ± 1·2</td>
<td>85·4 ± 7·1</td>
<td>12·6/12·6</td>
</tr>
<tr>
<td>22</td>
<td>7</td>
<td>0·30 ± 0·02</td>
<td>10·8 ± 3·2</td>
<td>29·7 ± 2·7</td>
<td>12·6/12·6</td>
<td>12·6/12·6</td>
</tr>
<tr>
<td>1 day</td>
<td>4</td>
<td>0·23 ± 0·05</td>
<td>0·8 ± 0·2</td>
<td>12·4 ± 2·7</td>
<td>—</td>
<td>—</td>
</tr>
</tbody>
</table>

Values are mean ± 1 s.e.m.

Experiment 2: steroid treatment of intact pregnant rats

Intact rats at Day 18 of pregnancy were treated s.c. at 09:00 and 16:00 h with 0·5 or 5 µg oestradiol benzoate/kg or 10 mg progesterone/kg. The steroids had no effect on cervical extensibility when measured on Day 19 (Table 2).
Table 2. Effect of oestradiol benzoate or progesterone given twice on Day 18 of pregnancy on the properties of the rat cervix (extensibility (10³ K/l₀) and inner circumference (l₀)) on Day 19

<table>
<thead>
<tr>
<th>Treatment</th>
<th>No. of rats</th>
<th>Wet wt (g)</th>
<th>Extensibility (min⁻¹)</th>
<th>Inner circumference (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>5</td>
<td>0.21 ± 0.03</td>
<td>7.5 ± 0.7</td>
<td>20.8 ± 2.5</td>
</tr>
<tr>
<td>Oestradiol benzoate</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0-5 µg/kg</td>
<td>8</td>
<td>0.19 ± 0.01</td>
<td>8.6 ± 3.3</td>
<td>20.1 ± 1.3</td>
</tr>
<tr>
<td>5-0 µg/kg</td>
<td>6</td>
<td>0.18 ± 0.01</td>
<td>5.9 ± 0.7</td>
<td>19.0 ± 0.7</td>
</tr>
<tr>
<td>Progesterone</td>
<td>7</td>
<td>0.22 ± 0.01</td>
<td>5.9 ± 1.1</td>
<td>18.2 ± 2.5</td>
</tr>
</tbody>
</table>

Values are mean ± 1 s.e.m.

Experiment 3: Ovariectomy and hormone treatment

Group 1. Following bilateral ovariectomy of rats on Day 16 of pregnancy most fetuses were alive in rats killed on Day 18 (Group 1a) but many were dead or resorbed in rats killed on Day 20 (Group 1b; Table 3). By Day 20 some fetuses had been expelled while others were lodged at the utero-cervical junction. Cervical wet weight and inner circumference had increased in these rats by Day 20 compared to Day 16 (Table 1) but extensibility had decreased to values seen in non-pregnant rats.

Table 3. Effect of bilateral ovariectomy on Day 16 of pregnancy with and without steroid treatment and relaxin (see text) on the properties of the rat cervix (extensibility (10³ K/l₀) and inner circumference (l₀)) and products of conception

<table>
<thead>
<tr>
<th>Group</th>
<th>Day killed</th>
<th>No. of rats</th>
<th>Wet wt (g)</th>
<th>Extensibility (min⁻¹)</th>
<th>Inner circumference (mm)</th>
<th>Wt (g)</th>
<th>Mean no. of fetuses alive/total</th>
</tr>
</thead>
<tbody>
<tr>
<td>1a. Ovariectomy only</td>
<td>18</td>
<td>9</td>
<td>0.16 ± 0.01a</td>
<td>3.9 ± 0.8a</td>
<td>16.3 ± 0.8a</td>
<td>24.5 ± 0.7</td>
<td>12.5/12.8a</td>
</tr>
<tr>
<td>1b. Ovariectomy only</td>
<td>20</td>
<td>7</td>
<td>0.28 ± 0.02a,b</td>
<td>1.0 ± 0.2a,b</td>
<td>29.9 ± 1.3a,b</td>
<td>21.9 ± 3.2a</td>
<td>3.1a,b/6.3a,b</td>
</tr>
<tr>
<td>2a. Steroids</td>
<td>19</td>
<td>7</td>
<td>0.11 ± 0.01c</td>
<td>2.2 ± 0.3c-f</td>
<td>12.5 ± 1.3c,d,e</td>
<td>31.7 ± 2.0</td>
<td>10.9/11.4</td>
</tr>
<tr>
<td>2b. Steroids</td>
<td>20</td>
<td>4</td>
<td>0.11 ± 0.01b</td>
<td>2.8 ± 0.3b</td>
<td>13.7 ± 1.7b</td>
<td>53.8 ± 1.3a</td>
<td>13.5b/13.8b</td>
</tr>
<tr>
<td>3. Progesterone withdrawal</td>
<td>19</td>
<td>8</td>
<td>0.13 ± 0.01b</td>
<td>2.7 ± 0.5</td>
<td>16.6 ± 0.7c</td>
<td>40.0 ± 1.7</td>
<td>14.5/14.5</td>
</tr>
<tr>
<td>4. Steroids + relaxin</td>
<td>19</td>
<td>7</td>
<td>0.12 ± 0.00d</td>
<td>3.3 ± 0.3c,d</td>
<td>17.9 ± 1.0d</td>
<td>35.1 ± 1.6</td>
<td>12.3/12.4</td>
</tr>
<tr>
<td>5. Steroids + relaxin</td>
<td>19</td>
<td>4</td>
<td>0.16 ± 0.01c,d</td>
<td>5.3 ± 0.8c,d</td>
<td>17.8 ± 1.6e</td>
<td>36.2 ± 1.8</td>
<td>13.0/13.0</td>
</tr>
</tbody>
</table>

For details of group treatments see text. Values are mean ± 1 s.e.m. Values within columns with same superscripts are significantly different (a, b, f 2P < 0.05; c, d, e 2P < 0.01).

Group 2. To replace ovarian steroid secretion, other groups of bilaterally ovariectomized rats were injected s.c. with 0.5 µg oestradiol benzoate/kg plus 10 mg progesterone/kg, once on Day 16 and twice daily on Days 17 and 18 (Group 2a) or Days 17 to 19 (Group 2b), and killed 1 day later (Table 3). This regimen allowed the pregnancies to continue with a normal increase in conceptus weights. However, cervical extensibility had only increased to a small extent, much less than the increase that normally occurs over this period of pregnancy (Table 1).

Group 3. Another group of bilaterally ovariectomized rats were treated s.c. with 0.5 µg oestradiol benzoate/kg once on Day 16 and twice daily on Days 17 and 18 but 10 mg progesterone/kg once on Day 16 and twice on Day 17 only (i.e. progesterone withdrawal). The rats
were killed on Day 19. In this group inner circumference showed a small but significant increase compared to that in Group 2a but extensibility had not increased (Table 3).

Groups 4 and 5. In addition to the steroid regimen of Group 2a, animals in Group 4 received s.c. injections of 1000 units relaxin (NIH-R-P1)/kg twice on Day 18 only, while those in Group 5 received 0.5 mg relaxin (CMa1-18-28AAE)/kg once on Day 16 and twice on each of Days 17 and 18. In both groups there was increased cervical extensibility on Day 19 and small increases in inner circumference (Table 3).

Discussion

The simple method described in this paper permits measurement of the large increase in extensibility and inner circumference of the rat cervix that occurs during pregnancy, in preparation for the need to dilate at term, and the changes in these produced by hormone manipulation. The increases observed in untreated pregnant rats in the present results confirm the observations of Harkness & Nightingale (1962) and Rundgren (1974).

The cervix of the late pregnant rat contains smooth muscle which is capable of responding to prostaglandins, oxytocin and neurotransmitters, but on Day 22 maximal responses of spirally cut cervical strips were small relative to those of the uterine horn (Hollingsworth & Isherwood, 1978, 1979), perhaps due to the lower proportion of smooth muscle in the cervix (Harkness & Harkness, 1959). It is therefore reasonable to account for changes in the extensibility of the cervix mainly in terms of changes in the connective tissue.

There is only limited information on the possible hormonal control of the increase in cervical extensibility during late pregnancy in the rat (see ‘Introduction’). The present results and those of others (Csapo & Wiest, 1969; Català & Deis, 1973; Buckle & Nathanielsz, 1975) show that parturition after ovariectomy in late pregnancy in the rat is protracted, fetal and neonatal deaths occur and often fetuses are lodged in a cervical canal. This could be due to a lack of an ovarian hormone to provide the terminal increase in cervical extensibility. Sufficient cervical dilatation could still occur to allow fetal expulsion as fetal head circumference would be less than at term and the stresses of the fetal head would appear to be exerted for a longer period of time. Uterine motility, as measured by intrauterine balloons in conscious rats, was considerably greater in ovariectomized than control pregnant animals (M. Hollingsworth & C. N. M. Isherwood, unpublished).

The ovarian hormone producing increased cervical extensibility could be an oestrogen. Oestrogens increased cervical weight and inner circumference in the non-pregnant rat but produced only small increases in cervical extensibility (Kroc et al., 1959; Cullen & Harkness, 1960; Zarrow & Yochim, 1961). The changes in the cervix described by Kroc et al. (1959) and Zarrow & Yochim (1961) are probably mainly reflections of increased tissue mass rather than greater extensibility. Oestradiol administered in large doses to women at term produced some cervical softening as judged clinically (Pinto, Fisch, Schwarz & Montiori, 1964; Gordon & Calder, 1977). In the present work, oestradiol benzoate and progesterone had no effect on cervical extensibility when given to intact rats but the combination of progesterone and oestradiol benzoate produced a small increase when given to ovariectomized rats. The dosages of progesterone and oestradiol benzoate used are similar to those which will, respectively, return plasma progesterone concentrations to normal values following ovariectomy (Csapo & Wiest, 1969) and produce normal delivery following ovariectomy near term (Català & Deis, 1973). Also, the increases in extensibility during pregnancy described here and by Harkness & Nightingale (1962) and Rundgren (1974) precede the rise in plasma oestradiol concentrations near term (Shaikh, 1971).

Progesterone is able to increase slightly cervical weight and inner circumference above that of oestrogen-treated rats but does not increase extensibility (Kroc et al., 1959; Cullen &
Progesterone concentrations in the late pregnant rat from about Day 17 or 18 (Csapo & Wiest, 1969; Buckle & Nathanielisz, 1975) increased cervical extensibility could be a direct consequence of progesterone withdrawal. However, cervical extensibility decreases following ovariectomy when plasma progesterone concentrations will have fallen (Csapo & Wiest, 1969) and extensibility could not be significantly increased with the progesterone-withdrawal treatment used in the present study. Fitzpatrick (1977a, b) has suggested that increased cervical softening can only be produced in sheep with dexamethasone or prostaglandin F-2a or in the goat with cloprostenol when there are low progesterone concentrations or a low progesterone/oestrogen ratio. Progesterone may have an inhibitory action on the cervix, preventing other hormones from increasing cervical extensibility. Other methods of producing progesterone withdrawal are necessary to test this further.

Another hormone synthesized and stored in rat ovaries is relaxin (Anderson & Long, 1978). Serum relaxin concentrations in pregnant rats, as measured by radioimmunoassay, reach a peak about Day 15 and then slowly decline (O’Byrne & Steinetz, 1976). This peak precedes the rise in cervical extensibility in rats and there is some correspondence between serum relaxin concentrations and increased cervical size in the pregnant hamster (O’Byrne, Sawyer, Butler & Steinetz, 1976). Ovariectomy would remove the source of relaxin in the rat and this could explain the minimal increase in cervical extensibility in the ovariectomized, steroid-treated group. Treatment with relaxin increased cervical extensibility and further experiments are necessary, using different doses and schedules of relaxin, to determine whether extensibilities equivalent to those seen at term could be produced in ovariectomized, steroid-treated rats.

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