Sex differences in plasma corticosterone in mouse fetuses are mediated by differential placental transport from the mother and eliminated by maternal adrenalectomy or stress

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The effect of changes in maternal corticosterone concentrations, induced by maternal stress, maternal adrenalectomy or both, on concentration of corticosterone in serum and in adrenals of mouse (Mus domesticus) fetuses was examined. Higher baseline serum corticosterone concentrations were found in female fetuses than in male fetuses; however, there was no sex difference in the content of corticosterone in adrenals collected from these fetuses. Sex differences were observed in the fetal response to changes in maternal concentrations of serum corticosterone resulting from stress (bright light and heat) or adrenalectomy, and both factors eliminated the sex difference in corticosterone in fetal serum. When females were injected i.p. with 14C-corticosterone on day 17 of pregnancy, significantly more 3H was recovered from the serum of female than of male fetuses 15 min after the injection, while more 1H was recovered from placenta of male fetuses. This finding suggests that the difference in serum corticosterone in male and female mouse fetuses is due to greater transport of corticosterone from maternal blood across the placenta of female than of male fetuses.

Introduction

The fetal pituitary-adrenal system starts functioning between days 17 and 18 of pregnancy in rats (Dapouy et al., 1973), and an important modulating role for corticosterone of maternal origin is well established (Yamakoshi et al., 1983). In rats the fetal adrenals hypertrophy between day 18 and day 22 of pregnancy after maternal adrenalectomy on day 14. This has been interpreted as a compensatory response of the fetal pituitary-adrenal axis to a fall in circulating glucocorticoids derived from an adrenalectomized mother; corticosterone, but not ACTH (which would be increased in mothers after adrenalectomy), crosses the placenta from the maternal to the fetal circulation (Zarrow et al., 1970; Wong and Burton, 1974; Dapouy et al., 1980). The presence of glucocorticoid receptors in the placenta has been reported in rats (Heller et al., 1981, 1983) and mice (Wong and Burton, 1974). Glucocorticoids may play a role in the inhibition of placent al progesterone secretion during mid-pregnancy (Heller and DeNicola, 1983). Most progesterone in the maternal circulation is secreted by the maternal ovaries (Pollini et al., 1981). The decrease in placental progesterone secretion that occurs at mid-pregnancy is associated with an induction of corticosterone metabolism by the placenta (Pepe and Albrecht, 1984), and, thus, a decrease in the passage of corticosteroids from the maternal to the fetal circulation. In mice, this occurs on day 11 (Montano et al., 1991), when there is a dramatic increase in corticosterone in maternal blood following a marked increase in placental estrogen secretion on days 9 and 10 of pregnancy (Barkey et al., 1979; Soares and Talamantes, 1982, 1983; F. S. vom Saal, unpublished). Androgens synthesised by the placenta are secreted into maternal and fetal compartments and are substrates for estrogen biosynthesis in the maternal ovary (Jackson and Albrecht, 1985; Gilbert et al., 1986) and possibly also in fetuses (vom Saal et al., 1992). The dramatic changes in maternal glucocorticoid concentrations beginning during mid-pregnancy could influence steroid hormone (estrogen, androgen, glucocorticoid) concentrations in the fetal circulation. In addition, it is possible that sex differences in placental binding or metabolism of corticosteroids could be due to serum differences in these steroids between male and female fetuses (Ward and Weiss, 1980; Wilke et al., 1982; vom Saal et al., 1990; Montano et al., 1991).

Changes in maternal glucocorticoid concentrations owing to maternal stress or maternal adrenalectomy also influence androgens and glucocorticoids concentrations in the fetal circulation (Ward and Weiss, 1980; vom Saal et al., 1990; Montano et al., 1991). We, therefore, examined the effects of changes in maternal concentrations of serum corticosterone induced by maternal stress, adrenalectomy, or both, on serum corticosterone concentrations and adrenal corticosterone content in male and female fetuses. We also examined the transport of [14C]corticosterone from the maternal circulation into male and female fetuses.

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Materials and Methods

Animals

CF-1 mice were housed in 18 cm × 28 cm × 15 cm polypropylene cages on Asept bedding. Mouse breeders in (Partula 50000) and water were available ad libitum. Animal rooms were maintained at 23 ± 2°C in a 12 h light:12 h dark cycle, with lights on at 12:00 h. The onset of the light phase of the cycle will be referred to as 00:00 h, and all times of day will be relative to the beginning of the light phase of the cycle. All work in the animal room during the dark phase of the cycle was conducted under a 25 W red light to which mice do not respond.

Mating

Adult CF-1 female mice were mated by being placed with males for 4 h beginning at 21:00 h (0.5 h before the onset of the light phase of the light:dark cycle). Mating was confirmed by the presence of a copulatory plug and noted animals were considered to have been fertilized at the onset of the light phase at 00:00 h, which was thus labelled as the beginning (day 0) of pregnancy. Mated females were housed three per cage until assigned to a treatment group.

Surgery procedure

Randomly selected pregnant females were stressed using procedures similar to those described by vom Saal et al. (1990). Pregnant females were placed in Plexiglas mouse restraining bars (9 cm × 6.3 cm × 5 cm) under a bank of 150 W flood lights (3700 K at 38° C). Females were stressed twice a day at 08:00 h and 20:00 h beginning at 20:00 h on day 12 of pregnancy. Each stress session lasted 45 min. Control pregnant females were left undisturbed in their cages.

Surgery and collection of blood samples

Adrenoreceptors were performed via the dorsal approach at the beginning of day 11 of pregnancy using inhalation of methoxyflurane (Metofane; Pitman-Moore, NJ) as an anesthetic. All adrenoreceptorized females were housed singly and given access ad libitum to a solution of 1% saline (to compensate for the loss of adrenergic) and 0.5% deconex (for palatability) in tap water.

Pregnant females from all groups were housed individually on day 11 of pregnancy. On different days of pregnancy, females were decapitated, and blood was collected in test tubes within 15 s. All fetuses were removed from the uterine horns, and fetal blood was collected in heparinized microtubes by aspiration after fetal decapitation (within 2 mm of maternal decapitation). Fetal sex was determined by examining the length of the anogenital space, which unambiguously distinguishes males from females at the times examined in these studies. Blood from fetuses of each sex within a litter was pooled to control for litter effects. Fetuses were placed in ice until their adrenal were collected. The adrenals were frozen at -70°C immediately after removal. Blood was centrifuged and plasma frozen at -70°C until assayed.

After decapitation and removal of fetuses, adrenalectomized females were autopsied and tissue around the area of the excised adrenal was collected and frozen. The tissue was excised, weighed, and reserved for the presence of corticosterone. There was no indication of residual corticosterone-secreting adrenal tissue in any of the animals.

A preliminary study for residual corticosterone-secreting adrenal tissue after adrenalectomy

A preliminary experiment was performed to determine whether residual adrenal cortical tissue was present after adrenalectomy. Virgin rather than pregnant females were used as fetuses serve as a non-natural source of corticosterone in plasma of pregnant females. Ten virgin female mice were adrenalectomized at 12:00 h (just before the end of the light phase of the cycle), and blood was collected by decapsulation at either 24 h or 48 h (n = 5 females per group) after adrenalectomy (for examination of concentrations of corticosterone in plasma). Tissue samples were also collected from around the dorsal aspect of the kidney and examined for the presence of corticosterone.

At 24 h after adrenalectomy, two of five virgin females still exhibited concentrations of corticosterone in plasma (25 µg in 100 ml and 20 µg in 100 ml) that were similar to the peak values observed in intact virgin females killed at the same time of day (Montano et al., 1994). The concentrations of corticosterone in plasma of the other three females were 30% lower than values observed in intact females. By 48 h after adrenalectomy, no concentrations of corticosterone in plasma had fallen to 1:92 ± 0.03 µg/ml, 92% lower than values observed in intact virgin female mice killed at the same time of day. Corticosterone was not detected in tissue samples collected along the anterior border of the kidney in any of the females used in this study. Taken together, these findings suggest that all corticosterone-secreting adrenal tissue was removed during the adrenalectomy procedure.

Extraction of steroids

Corticosterone was extracted twice with 2 ml of an ethylacetate:chloroform (90:20; v:v) mixture, dried under nitrogen and reconstituted in 140 µl buffer. It was necessary to dilute maternal plasma (1:40) in assay buffer to obtain values that fell within the range of the assay. Steroids were extracted from fetal adrenals in the same manner, except that there was a 30 h incubation after addition of the extraction mixture. Recovery was monitored by addition of 100 µg (Lot. 97, Sigma Chemical Co., St. Louis, MO) corticosterone (888.4 µg/mg) and New England Nuclear, Boston, MA) to control samples before extraction. A 20 µl aliquot for recovery and drug standard stock aliquots for steroids radioimmunoassay were withdrawn after reconstitution. Recovery was routinely greater than 90%.

Corticosterone radioimmunoassay

Corticosterone radioimmunoassay was performed as described in detail by Montano et al. (1994). Briefly, 1,26-3H corticoster-
one (0.5G C mmol-1) was obtained from New England Nuclear (Boston, MA) and unlabeled corticosterone was obtained from Steraloids, Inc. (Wilton, NH). Corticosterone antiserum (rabbit) was obtained from Gordon/Novooxides (P.Collins, CO). Duplicate aliquots of 200 μl of extracted steroid in buffer were incubated for 4 h at 24°C with 18,000 c.p.m. (33 pg) of [1]H)corticosterone and antiserum to corticosterone. Free and bound steroids were separated by second antibody addition (diluted 1:1) goat antirabbit; Radiassay Systems Laboratories, Inc. (Canons, CA). Tubes were incubated overnight at 4°C and then centrifuged at 1500 g for 1 h. The supernatant was decanted, and the precipitate (representing the bound portion) was counted in a liquid scintillation counter (Beckman LS 5801, efficiency for tritium: 56%) after addition of 9 ml Scintiverse (I/P). The range of the standard curve for corticosterone was 0–3000 pg per tube. The blank of the assay was indistinguishable from baseline values. The inter- and intra-assay coefficients of variation were 11% and 3%, respectively.

Statistical analysis

Serum and tissue corticosterone concentrations were compared by analysis of variance or analysis of covariance using the Statistical Analysis System (SAS, General Linear Model). Planned comparisons were made using the LS means test (SAS).

Experimental Details and Results

Experiment 1. Effect of adrenalectomy on day 11 on corticosterone concentrations in plasma of pregnant females

The objective of this experiment was to assess when the adrenals of fetuses begin to secrete corticosterone Examining corticosterone concentrations in maternal blood after adrenalectomy on day 11 of pregnancy. Before day 16 of pregnancy, too little blood was collected from fetuses to assess directly the onset of secretion of corticosterone in fetuses.

Method. Pregnant females were adrenalectomized at 00:00 h on day 11 of pregnancy. Blood was collected for determination of concentrations of corticosterone in plasma at 00:00 h just after the onset of the eight phase of the light:dark cycle on days 14–16 of pregnancy. Blood was also collected from intact pregnant females at corresponding times. Seventy-five pregnant females were used.

Results. Analysis of variance performed on maternal serum corticosterone revealed a significant interaction between day and treatment (P < 0.001). Females adrenalectomized on day 11 of pregnancy showed serum corticosterone concentrations on day 14 that were 3% of the concentrations (5.0 ± 0.8 pg/100 μl) observed in intact pregnant females (553 ± 7 pg/100 μl). An inverse in plasma corticosterone was observed on day 15, followed by concentrations on day 16 that were not significantly different from those of intact pregnant females (Fig. 1). Adrenalectomized females had significantly higher concentrations of corticosterone in plasma than did intact females on day 17 (P < 0.001) and day 18 (P < 0.005).

Fig. 1. Concentrations of corticosterone in plasma (mean ± SEM) in (1) intact and (•) adrenalectomized (ADX) on day 11 of pregnancy females (n = 9) on days 14–16 of pregnancy. Number of samples assayed for successive time points were: intact n = 7, 13, and 12; adrenalectomized n = 5, 5, 11, and 9. Significant difference between intact and adrenalectomized females (P < 0.001). *Significant difference between intact and adrenalectomized females (P < 0.001).

Experiment 2: effect of maternal adrenalectomy on day 11 on stress on corticosterone in maternal and fetal serum and fetal adrenal glands

In this experiment, the fetal response to maternal adrenalectomy, stress and a combination of adrenalectomy and stress on day 18 of pregnancy were determined. Eighteen-day-old fetuses were used as the previous experiment showed evidence of hypersensitivity by the fetal adrenals on day 18 following maternal adrenalectomy on day 11, and a much greater volume of plasma can be obtained from fetuses on day 18 than on day 17 of pregnancy.

Methods. Pregnant females were adrenalectomized on day 11. Half of the adrenalectomized females, and half of the group of intact females were subjected to stress twice each day: starting at 20:00 h on day 13. This was to allow sufficient time for recovery after adrenalectomy. A group of adrenalectomized and intact females was left undisturbed until the time of blood and tissue collection (referred to as control-adrenalectomized and control-intact females, respectively).

Blood was collected from stressed and control animals at 00:00 h on day 18 1 h after the onset of the last stress session, which began at 20:00 h on day 17 during the dark phase of the light:dark cycle. We previously observed that concentrations of corticosterone in plasma were relatively low at 00:00 h on day 18 in intact, non-stimulated mouse fetuses (Meninano et al., 1993). Thirty-five fetuses were used.

Results. Control-adrenalectomized mothers again showed higher concentrations of corticosterone relative to control-intact mothers, although the difference was not statistically significant. As expected, there was no significant difference in plasma corticosterone between stressed-adrenalectomized and control-adrenalectomized pregnant females. However, there tended to be an increase (P = 0.07) in serum corticosterone in stressed-intact mothers relative to control-intact mothers 4 h
Fig. 2. Concentrations of corticosterone in plasma (mean ± SEM) in intact and adrenalectomized (on day 11) pregnant females at 00:00 h on day 18 of pregnancy. Stress (X) stressed twice a day starting on day 13. Control (□) left undisturbed until collection of blood samples. Intact control, n = 6; intact stress, n = 13; adrenalectomized (ADX) control, n = 6; adrenalectomized stress, n = 10.

Fig. 3. Concentrations of corticosterone in plasma (mean ± SEM) on day 18 of pregnancy at 00:00 h in male and female fetuses carried by intact and adrenalectomized (ADX) (on day 11) mothers. Stress mothers were stressed twice a day starting on day 13. Control mothers were left undisturbed until sample collection. The following number of samples were assayed (representing pooled sera from fetuses of the same sex in each litter): male fetuses (♂), intact control, n = 7; intact stress, n = 13; adrenalectomized control, n = 9; adrenalectomized stress, n = 8; female fetuses (♀), intact control, n = 6; intact stress, n = 11; adrenalectomized control, n = 7; adrenalectomized stress, n = 8. *Significant difference between male and female fetuses (P < 0.01).

After the last stress session (Fig. 2), which was consistent with previous findings (Montano et al., 1993).

In fetuses from intact, control mothers, females had significantly (P < 0.01) higher concentrations of corticosterone in plasma than did males (P < 0.01). Figure 3. However, no sex difference in fetal adrenal corticosterone content was observed in these fetuses (Fig. 4). Maternal stress induced a decrease in serum corticosterone concentrations relative to control fetuses from intact mothers only in female fetuses: corticosterone decreased (P < 0.05) to control male values (Fig. 3). In both male and female fetuses from intact mothers, only a slight but nonsignificant decrease in adrenal corticosterone content was observed as a result of maternal stress (Fig. 4).

Maternal adrenalectomy induced an increase in serum corticosterone concentrations in male fetuses (P < 0.05) but not in female fetuses. In contrast, there was a significant difference in adrenal corticosterone content in fetuses (regardless of sex) from intact versus adrenalectomized mothers (P < 0.05).

Experiment 3: Effect of adrenalectomy on day 18 of pregnancy on corticosterone in plasma of pregnant mothers and fetuses

The results of Exp1 suggested that fetuses begin secreting corticosterone as early as days 14–15 of pregnancy. Furthermore, the high corticosterone concentrations in maternal serum on day 17 and 18 following adrenalectomy on day 11 appeared to reflect hypersecretion of corticosterone by the fetal adrenals, since adrenal corticosterone content was high in both male and female fetuses carried by adrenalectomized mothers (Exp2). The objective of this experiment was to determine whether hypersecretion by the fetal adrenals on day 18, which was observed after maternal adrenalectomy on day 11 in Exp1, required 0–7 days or would also be observed within 1–2 days after maternal adrenalectomy on day 16.

Material: Pregnant females were adrenalectomized on day 16 of pregnancy at 00:00 h. Separate groups of females were killed at 00:00 h on day 17 and day 18, and blood was collected from both the pregnant females and their fetuses.
Fig. 5. Concentrations of corticosterone in plasma (mean ± SEM) in intact (C) and adrenalectomized (ADX) on day 10 of pregnancy. Control, left undisturbed until blood collection. The following number of samples were analyzed: day 17 intact, n = 8; adrenalectomized, n = 7; day 18 intact, n = 6; adrenalectomized (ADX), n = 4. Significant difference between intact and adrenalectomized pregnant females (P < 0.05).

Results. Analysis of variance performed on concentrations of corticosterone in plasma of mothers revealed a significant interaction between day and treatment (P < 0.01; Fig. 5). After adrenalectomy on day 10, significantly higher concentrations of corticosterone in plasma were observed on day 17 of pregnancy of intact (control) pregnant females relative to adrenalectomized pregnant females (P = 0.03). Corticosterone plasma tended to be higher on day 18 in adrenalectomized females relative to control females (P = 0.07).

Analysis of serum corticosterone concentrations on day 17 and 18 in fetuses revealed significant main effects of day and treatment (day: P < 0.01; treatment: P < 0.05). Overall, concentrations of corticosterone in plasma were lower on day 18 than on day 17 (Fig. 6). Irrespective of day of pregnancy, male fetuses from adrenalectomized mothers showed significantly higher plasma corticosterone levels compared to intact male fetuses from intact mothers, whereas there was no significant difference in concentrations of corticosterone in plasma of female fetuses from adrenalectomized versus intact mothers. Concentrations of corticosterone in plasma of female fetuses from intact mothers were 34% higher than concentrations of corticosterone in plasma of female fetuses on day 18. The difference was not statistically significant.

Experiment 4: transport of \( ^{3}H \) corticosterone across the placenta from maternal to fetal blood.

In Exp 4 we observed that female fetuses had higher concentrations of corticosterone in plasma than did male, whereas the adrenal cortex of corticosterone was virtually identical in these males and females. Changes in corticosterone concentrations of maternal plasma induced by maternal adrenalectomy, stress or a combination of both) led to markedly different effects on corticosterone in plasma of male and female fetuses. However, there was no difference between male and female fetuses in adrenal corticosterone content in response to these manipulations. These findings suggest that the sex difference in corticosterone of plasma in fetuses observed on day 18, as well as the sex difference in response to maternal stress and adrenocorticotropin, is not due to a sex difference in the functioning of the brain—hypothalamic—adrenal axis during fetal life. We therefore examined the possibility that there are differences in placental transport and placental binding of corticosterone between male and female fetuses.

Methods. We injected 3 µCi (17 ng) of \( ^{3}H \) corticosterone (301.6 Ci mmol\(^{-1} \)) New England Nuclear, Boston, MA) in 100 µal saline (with 2% ethanol) into the tail vein of female mice at 00:00 h on day 18 of pregnancy. In a previous study we observed a significant difference in concentrations of corticosterone in plasma between male and female fetuses at this time (Morris et al., 1993). Pregnant females were decapitated, and trunk blood was collected 15–20 min after injection of \( ^{3}H \) corticosterone into the tail vein and within 35 s of the cage being touched. In a preliminary study the highest amount of radioactivity was found in fetal plasma at 15–20 min (relative to 00 or 12) after injection of \( ^{3}H \) corticosterone into pregnant females.

Sixty fetuses (25 males and 35 females) from six litters were examined. All fetuses were removed from the uterine horns. Male and female fetuses were identified by examining anogenital distance, and blood was collected from individual fetuses in heparinized microcentrifuge tubes. After fetal decapitation within 2 min of maternal decapitation, the placenta from each fetus was weighed and placed in a 20 ml scintillation vial in an ice bath. Blood from individual fetuses was centrifuged at 1000 g for 30 min, and duplicate 10 µl aliquots were placed in 5 ml scintillation vials. Triplicate aliquots of the injection solution were also placed in scintillation vials and scintillation fluid was added for reference counting. Plasma was solubilized in 1 ml of Scholes-\( ^{3}H \) (Packard) at 80°C for 1 h. After addition of scintillation fluid vials containing plasma and placental samples, the
samples (including reference visits) were counted in a Beckman LS-5861 counter. Parameters were programmed in the liquid scintillation counter so that quench corrections were performed automatically for each sample.

Since metabolites of $[^{3}H]$ corticosterone were not recovered, the d.p.m. recovered from plasma and plasma will be referred to as $[^{3}H]$. Pearson correlation coefficients were calculated for $[^{3}H]$ present in maternal serum, plasma and fetal plasma. The correlation coefficients were also calculated for both plasmatic weight and the ratio of males and females within a litter and all of the above variables.

Results. There was an average of 11.7 ± 0.7 fetuses per litter, and number of male fetuses number of female fetuses was $0.71 ± 0.1$. Sex ratio was not significantly correlated with plasmatic weight, $[^{3}H]$ in plasma or $[^{3}H]$ in fetal blood. Male plasmatic weight was 10.8% heavier than female plasmatic weight: males = 849 ± 2.9 mg, females = 613 ± 2.8 mg; $P < 0.005$, whether analyzed by ANOVA or using sex ratio as the covariate.

Correlation analysis ($n = 70$) showed that plasmatic weight was the greatest predictor of $[^{3}H]$ in the plasma $r = 0.69, P < 0.001$. When analyzed by analysis of covariance using sex ratio as the covariate, the ratio of $[^{3}H]$ recovered from male plasma ($15,500$ d.p.m.) was $44%$ higher than from female plasma ($13,374$ d.p.m.; $P = 0.001$). Significant negative correlation of corticosterone in plasma and plasmatic weight, although when analyzed by ANOVA this sex difference was not as great ($P = 0.12$).

Radioactivity in fetal plasma (males = 118.571 d.p.m. in 10 µl plasma, females = 107.691 d.p.m. in 10 µl plasma) was significantly correlated with $[^{3}H]$ in maternal plasma (males = 7.9%, females = 0.001). The ratio of $[^{3}H]$ in fetal plasma $[^{3}H]$ in maternal plasma was significantly (males = 0.005) higher for females than for male litters suggesting a greater transport of corticosterone from maternal plasma into female than into male fetuses.

Experiment 5: Adrenal weight and content of corticosterone in adult female mice

Sex differences in adrenal weight and concentrations of corticosterone in plasma have been reported for mice, but not all, strains of mice (Shire, 1974). Since a sex difference in adrenal corticosterone content was not found in fetuses, adrenal weight and corticosterone content were compared in adult, virgin male and female C3H mice to examine the possibility that this lack of mice does not show a sex difference in adrenal corticosterone content even in adulthood.

Methods. Animals of the same sex were housed four per cage from weaning until the day before being killed (when 90 days old), at which time each animal was individually housed. Animals were killed by cervical dislocation at 12.00 (before peak values are observed in adult, virgin mice; Montano et al., 1991), and the adrenal glands were removed and weighed (24 animals were used).

Results. Analysis of variance revealed significantly (by 5%) higher adrenal weights (mg per 100 g body weight) for paired adrenal glands in adult female mice than in adult male mice (females = 21.2 ± 2.5 vs. males = 13.6 ± 0.8; $P < 0.001$). There was also a significant sex difference (86% higher in females than in males) in adrenal corticosterone content (females = 110 ± 5 ng per adrenal; males = 59 ± 8 ng per adrenal; $P = 0.005$). Concentrations of corticosterone in plasma were 110% higher in females than in males (females = 16.9 ± 2.5 µg in 100 ml; males = 6.9 ± 1.0 µg in 100 ml; $P < 0.001$).

Discussion. In female mice adrenalized on day 11 of pregnancy, plasma corticosterone increased from very low concentrations on day 4 to concentrations observed in intact pregnancy females by day 16. It thus appears that by days 14-15 of pregnancy, secretion of corticosterone by the fetal adrenal begins and try day 16 can result in normal concentrations of corticosterone in maternal serum after maternal adrenalinization. The rise in maternal serum corticosterone after maternal adrenalinization in pregnant rats has been correlated with the onset of fetal brain-pulmonary-adrenal activity, which has been reported to occur in rats between days 17 and 18 (Dopay et al., 1975) or days 19 and 20 (Chabot et al., 1968). Day 17-18 in pregnancy in rats roughly corresponds to days 16-17 of pregnancy in mice (Schild et al., 1967, Raphael, 1969). The time of onset of fetal brain-pulmonary-adrenal activity suggested by our findings in mice is thus earlier than that observed in rats. The significantly higher concentrations of circulating corticosterone in adrenalized pregnant female mice relative to intact females on days 17 and 18 was unexpected and was not evident in previous studies with rats (Milovanovic et al., 1973, Dopay et al., 1975, Chabot et al., 1968).

Adrenal content of corticosterone on day 18 of pregnancy was high in both male and female fetuses carried by mothers adrenalized on day 11. This finding is consistent with the predictions that there should be an increase in the fetal secretion of corticosterone releasing factor (CRF; Chabot et al., 1968), ACTH and corticosterone following a decrease in the fetal blood of corticosterone of maternal origin, since corticosterone, but not ACTH, can cross the placenta from the maternal to the fetal circulation (Dopay et al., 1980). Steroids are not stored within cells in placenta that produce them. The rate of secretion of steroid hormones relates the rate of synthesis and is correlated with organ content (Sauer and Talalansky, 1983). It thus appears that the increase in adrenal corticosterone content at the single point measured in fetuses reflected an increase in the rate of synthesis and secretion of corticosterone by the fetal adrenal glands in response to maternal adrenalinization. A sex difference was not observed in adrenal corticosterone content of fetuses carried by either intact or adrenalinized on day 11 (d) inoculated. This finding suggests that there is no sex difference in fetal adrenal corticosterone synthesis or secretion. However, an increase in adrenal female plasma was observed on day 18 in male fetuses carried by mothers adrenalinized on either day 11 (Exp 2) or day 16 (Exp 3), whereas no change was observed in concentrations of corticosterone in plasma of female fetuses after maternal adrenalinization on either day. Maternal adrenalization, and the loss of corticosterone of maternal origin in the fetal blood, thus eliminated the difference in plasma corticosterone between male and female fetuses.
There was an increase in corticosterone in maternal plasma 4 h after the onset of maternal stress (Exp. 2). The increase in maternal corticosterone owing to stress led to a suppression of plasma corticosterone in female fetuses (at increased male values). This finding suggests a suppression of corticosterone secretion by the adrenal glands of female fetuses carried by stressed mothers, but adrenal corticosterone content was only slightly (and not significantly) lower in stressed female fetuses relative to control female fetuses. In addition, a change in plasma corticosterone was not observed on day 18 in male fetuses when blood was collected 4 h after the onset of maternal stress, although their adrenal corticosterone content was slightly decreased relative to control male fetuses. The basis for these findings is unknown and contrasts with previous findings obtained one day earlier on day 17 of pregnancy in mice (Montano et al., 1993).

The only sex difference observed in any of these studies was in plasma corticosterone in fetuses carried by mothers that were not stressed or adrenalectomized. We thus hypothesized that a sex difference in the passage of corticosterone from the maternal circulation through the placenta might mediate the sex difference in circulating corticosterone in fetuses, perhaps via a difference between male and female fetuses in placental binding of corticosterone. Between 35 and 20 min after injection of \( \text{ITI} \) corticosterone into pregnant females on day 18, significantly (15%) higher recovery of \( \text{ITI} \) in fetal female plasma was observed compared with male fetal plasma. This was accompanied by a 14% (\( P = 0.01 \)) higher degree of retention of \( \text{ITI} \) in male versus female placenta.

The above finding provides evidence that the higher corticosterone concentrations in plasma of female fetuses compared with male fetuses is due to a greater passage of corticosterone from the maternal blood across the placenta of female relative to male fetuses, as there does not appear to be a sex difference in the capacity of the fetal adrenal glands to synthesize corticosterone. Total \( \text{ITI} \) content in placenta and fetal blood was not examined and \( \text{ITI} \) labelled metabolites were not identified; the possibility that there is a sex difference in placental or fetal corticosterone metabolism, or in both, cannot therefore be excluded. In contrast to our findings in male and female fetuses, adult, virgin female CF-1 mice had significantly higher adrenal corticosterone content (and adrenal gland size) than did adult males.

On day 18 of pregnancy in CF-1 mice, there are sex differences in circulating testosterone (male fetuses have higher testosterone than do female fetuses) and estradiol (male fetuses have lower concentrations than do female fetuses; von Stall et al., 1990). Sex differences in the plasma concentrations of gonadal steroids during fetal life may play a role in mediating the sex difference in placental transport of corticosterone from the maternal to the fetal circulation.

Corticosterone-binding globulin is another factor that could play a role in mediating the sex difference in plasma corticosterone levels in females, as well as in the endocrine response of male versus female fetuses to changes in maternal plasma corticosterone concentrations. This protein binds with high affinity to glucocorticoids and affects the clearance rate and the bioavailability of glucocorticoids during development. Sex differences were observed in changes in plasma corticosterone in response to maternal adrenocortical and maternal stress only during periods where there were sex differences in control (baseline) plasma corticosterone concentrations. It remains to be determined whether a sex difference in corticosterone-binding globulin concentrations in fetal plasma plays a role.

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