Growth of suckling rats after treatment with dexamethasone or cortisol

Implications for steroid therapy in human infants

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De Souza, S. W., and Adlard, B. P. F. (1973). Archives of Disease in Childhood, 48, 519. Growth of suckling rats after treatment with dexamethasone or cortisol: implications for steroid therapy in human infants. Immature rats were treated with a single dose of dexamethasone (1 mg/kg or 20 mg/kg body weight). At both dose levels the subsequent growth of whole body, brain, and thymus was markedly impaired. Cortisol had smaller but similar effects. These results are discussed in the context of possible glucocorticoid therapy in newborn infants.

Methods

Animals. Rats of a black- and white-hooded strain were used. At birth, litters were reduced to either 6 (dexamethasone experiment, all males where possible) or 10 (cortisol experiment, equal numbers of each sex) animals.

Steroids. Dexamethasone phosphate (sodium salt) was a gift from Merke, Sharp and Dohme Ltd., Hoddesdon, Herts. Cortisol hemisuccinate (sodium salt) was purchased from Sigma Chemical Co., Kingston upon Thames, Surrey.

Brain dissection. Brain was separated from spinal cord at the foramen magnum. Cerebellum was routinely removed and weighed separately. The remainder of the brain is referred to as 'forebrain and brainstem'. The term 'whole brain' includes cerebellum.

Dexamethasone experiment. Rats in each litter were paired according to body weight and sex. Each member of a pair received, at 5 days of age, one dose of either dexamethasone phosphate dissolved in water, or an equal volume of isotonic glucose. (The quantity of glucose given was negligible compared with circulating amounts.) The dose of steroid was 1 mg/kg in 7 litters and 20 mg/kg in 5 litters. Two animals (one dexamethasone-treated rat together with its matched glucose-injected control) were killed from each litter at ages 7, 10, and 21 days.

Cortisol experiment. 6 litters of 10 animals each were used. At 4 days of age 1 animal of each sex was killed from each litter. The remaining 8 animals were randomly assigned to one of 4 treatment groups, each consisting of 1 member of each sex. Three groups received cortisol hemisuccinate (1 mg/kg, 5 mg/kg, 50

There is experimental evidence suggesting that glucocorticoids may be useful in the prevention of hyaline membrane disease (DeLemos et al., 1970; Kotas et al., 1971). Large doses of glucocorticoids are, however, well known to cause atrophy of lymphoid tissue and impair brain growth (Howard, 1965; Schapiro, 1965) in immature animals. Thus, before glucocorticoids are used in clinical practice, some consideration must be given to the possible harmful effects of these drugs on the developing organism (Reynolds, 1971). In addition, it has been suggested (Howard, 1968) that repeated stress may interfere with brain growth and cell division through increased adrenocortical secretion and raised plasma corticosteroid levels.

In the work reported here, dexamethasone was administered to developing rats in a single dose (1 mg/kg) comparable with that used clinically in infants, and in a higher dose (20 mg/kg) which has been found necessary to prevent the occurrence of cerebral oedema in asphyxiated immature rats (De Souza and Dobbing, 1973). Immature rats were also treated daily with cortisol either in a high dose (5 or 50 mg/kg) or in an amount (1 mg/kg) within the range which might be secreted in a stressed infant (Bertrand et al., 1963; Aarskog, 1965; Alleyne and Young, 1967).

Rats aged 4 or 5 days were used, since at this age rat brain is at a stage of development comparable with that of the human baby brain towards the latter part of the last trimester of pregnancy (Dobbing, 1970).
mg/kg body wt, intraperitoneally) dissolved in isotonic saline daily between 4 to 8 days of age. One group (controls) received the same volume of isotonic saline for the same period. All animals were killed at 9 days of age, 24 hours after the last injection.

DNA determination. Total brain DNA, a measure of total numbers of cells, was determined according to Zamenhof et al. (1964).

Results

Dexamethasone at both doses used resulted in an acute reduction in growth of body, brain, and thymus to 7 days of age (Table I). Thymus was markedly more affected than whole body, while deficits in cerebellar weight were greater than those of the remainder of the brain. The extent of growth retardation in 10-day-old dexamethasone-treated rats, previously reported in preliminary form (Adlard and De Souza, 1973), was similar to that of 7-day-old animals.

Ultimately, some recovery of growth appeared to occur in animals given the lower dose of dexamethasone (1 mg/kg) since weight deficits at 21 days of age were generally less than those at 7 days (Table I). However, such recovery was not evident for the high dose (20 mg/kg) of dexamethasone.

Deficits in brain weight in dexamethasone-treated animals at 10 and 21 days of age were associated with greater reductions in total brain DNA.

### TABLE I

**Body, brain, and thymus weight in 7-day-old and 21-day-old rats after one dose of dexamethasone at 5 days of age**

<table>
<thead>
<tr>
<th>Age (dy)</th>
<th>Dexamethasone (1 mg/kg)</th>
<th>Dexamethasone (20 mg/kg)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Control</td>
<td>Experimental</td>
</tr>
<tr>
<td>Body weight (g)</td>
<td>7</td>
<td>16 ± 2</td>
</tr>
<tr>
<td></td>
<td>21</td>
<td>49 ± 5</td>
</tr>
<tr>
<td>Forebrain + brainstem weight (mg)</td>
<td>7</td>
<td>616 ± 46</td>
</tr>
<tr>
<td></td>
<td>21</td>
<td>1147 ± 50</td>
</tr>
<tr>
<td>Cerebellum weight (mg)</td>
<td>7</td>
<td>46 ± 7</td>
</tr>
<tr>
<td></td>
<td>21</td>
<td>174 ± 10</td>
</tr>
<tr>
<td>Thymus weight (mg)</td>
<td>7</td>
<td>49 ± 10</td>
</tr>
<tr>
<td></td>
<td>21</td>
<td>195 ± 28</td>
</tr>
</tbody>
</table>

*Note: Results are given as mean ± SD. Each experimental (dexamethasone-treated) animal was paired with a littermate control of the same sex and similar body weight at 5 days of age. There were 4 to 7 pairs (each from a different litter) for each age and dose. Results were compared using the paired comparisons 't' test.

*P < 0.05, †P < 0.01, ‡P < 0.001, § not significant.

### TABLE II

**Whole brain DNA in 7-day-old, 10-day-old, and 21-day-old rats after one dose of dexamethasone at 5 days of age**

<table>
<thead>
<tr>
<th>Dose (mg/kg)</th>
<th>Age (dy)</th>
<th>μmol DNA-P per brain</th>
<th>Deficit in total DNA (%)</th>
<th>Deficit in brain weight (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Control</td>
<td>Experimental</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>7</td>
<td>1.62 ± 0.26</td>
<td>1.55 ± 0.11§</td>
<td>—</td>
</tr>
<tr>
<td></td>
<td>10</td>
<td>3.02 ± 0.51</td>
<td>2.63 ± 0.42†</td>
<td>13</td>
</tr>
<tr>
<td></td>
<td>21</td>
<td>4.23 ± 0.74</td>
<td>3.74 ± 0.47§</td>
<td>—</td>
</tr>
<tr>
<td></td>
<td>10</td>
<td>2.86 ± 0.29</td>
<td>2.18 ± 0.47∗</td>
<td>24</td>
</tr>
<tr>
<td></td>
<td>21</td>
<td>4.33 ± 0.63</td>
<td>3.86 ± 0.63†</td>
<td>11</td>
</tr>
</tbody>
</table>

*Note: Cerebellum was pooled with forebrain and brainstem for these analyses. Results are given as mean ± SD. Each experimental (dexamethasone-treated) animal was paired with a littermate control of the same sex and similar body weight at 5 days of age. There were 3 to 7 pairs (each from a different litter) for each age and dose. Results were compared using the paired comparisons 't' test. All experimental groups showed significant (P < 0.05) deficits in brain weight.

*P < 0.05, †P < 0.01, ‡P < 0.001, § not significant. DNA-P, DNA-phosphorus.
DNA (Table II), which were statistically significant except at 21 days in the lower dosed animals.

Repeated large daily doses of cortisol impaired growth of body, brain, and thymus (Table III), though this effect was clearly smaller than with comparable amounts of dexamethasone.

Discussion

The 5-day-old rat brain and the human brain in the perinatal period are in a similar phase of development (Dobbing, 1968, 1970). The adult number of neurons (with few exceptions) have already formed, glial cells have begun to appear, and the establishment of neuronal interconnexions has begun. In the human brain there is still appreciable glial cell formation during the first 18 postnatal months (Dobbing, 1970). Therefore, any adverse effects of steroid treatment in 4- and 5-day-old rats could, by implication, occur perinatally in human babies.

Deficits in brain weight in rats treated with dexamethasone and those treated with cortisol could be due partly to a reduction in cell numbers, though other developmental processes, including myelination (Howard, 1965), may also be altered. Other studies suggest that shortly after treatment of animals with cortisol cell mitosis is inhibited in brain (Balázs and Cotterrell, 1972) and thymus and lymph nodes (Stevens, Colessides, and Dougherty, 1965). In the present study cerebellar growth was more retarded than that of the remainder of the brain after glucocorticoid treatment, a type of selectivity also observed after undernutrition (Culley and Lineberger, 1968; Chase, Lindsley, and O’Brien, 1969) and hyperphenylalaninaemia (Chase and O’Brien, 1970) in developing rats. A possible explanation may be found in the very rapid rate of cell formation in the cerebellum during early postnatal life (Dickerson and Dobbing, 1966; Altman, 1970).

The rate of cortisol secretion is reported to be in the range of 1·0 to 1·5 mg/kg per 24 hr in newborn infants (Aarskog, 1965) and somewhat lower in older children (Bertrand et al., 1963). Neither body nor brain growth in the suckling rat were significantly affected by cortisol in a dose of 5 mg/kg per 24 hr; thymus growth was reduced at this dose, but not significantly so at 1 mg/kg per 24 hr (Table III). If it is reasonable to extrapolate between rat and man it seems likely that increased adrenocortical secretion induced by stress is unlikely to affect brain growth and that any effects on thymus growth would be relatively small.

However, the possible long-term effects of treating pregnant women or newborn babies with glucocorticoids must be considered. Cortisol-treated animals are not only growth retarded, but also show certain behavioural abnormalities (Howard and Granoff, 1968; Schapiro, Salas, and Vukovitch, 1970). In addition, retarded development of lymphoid tissue could have adverse effects on the body’s immune mechanisms.

It can be argued, however, that a direct extrapolation between rat and man is difficult, since human development takes place over a much longer period so that possible adverse effects of steroids are grossly exaggerated when studied in a rapidly growing species such as the rat. There is evidence also that the rat may be more sensitive than man to the thymolytic effects of glucocorticoids (Claman, 1972). Nevertheless, women who received glucocorticoids in pregnancy are known to have had
growth-retarded babies (Warrell and Taylor, 1968). It is clear that administered steroids have a profound and deleterious effect on growth, including that of the brain. This must be carefully borne in mind when their use is being considered in the newborn baby.

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