Plasma Corticosteroids in Healthy Twin Pairs

J. D. MAXWELL, J. A. BOYLE, W. R. GREIG, and W. W. BUCHANAN

From the University Department of Medicine, and Centre for Rheumatic Diseases, Royal Infirmary, Glasgow

Three different but related mechanisms influence corticosteroid secretion by the normal human adrenal cortex. Under basal conditions the concentration of plasma cortisol and the secretion of pituitary ACTH constitute a negative feedback system through which fluctuations in plasma cortisol concentration are minimized. Plasma cortisol levels are, however, higher in the morning than at night because of a diurnal rhythm in the rate of ACTH secretion. Both negative feedback and diurnal rhythm may be overcome by the raised steroid secretion mediated in response to stress which causes great increase in ACTH output, and results in plasma cortisol levels much above those found in basal conditions (James, Landon, and Fraser, 1968). A unified hypothesis to explain the performance of the adrenal cortex under both normal and stressful conditions has been proposed by Yates and Urquhart (1962) who suggested that plasma cortisol levels were regulated by a negative feedback control system which had a variable set-point.

In humans great variations in corticosteroid metabolism (for example the acceleration of corticosteroid removal in hyperthyroidism, or the impairment of corticosteroid removal in liver disease or myxoedema) are not accompanied by large changes in plasma corticosteroid concentration. This stability denotes effective control of plasma corticosteroid levels by correction of the adrenal cortical secretion rate for alterations in corticosteroid metabolism.

The effect of environmental factors on adrenal cortical activity—for example the increase in adrenal corticosteroid secretion and increase in plasma cortisol levels in response to stress, or the alteration in the diurnal rhythm by variation in the activity cycle or length of day—is well recognized. But there is very little information available on the possible contribution of genetic factors in the control of pituitary-adrenocortical activity, or of plasma cortisol levels in man. Great variations in plasma cortisol levels have been found at a given time of day in different individuals thought to be in a similar physiological state, while plasma cortisol levels are relatively constant in any one subject. This inter-person variation and intraperson constancy has not been adequately explained. It does not appear to be due to differences in age, sex, weight, or build, but genetic factors might be important.

Racial differences in urinary steroid excretion have been reported by several workers. Politzer and Tucker (1958) and Edozien (1960) found low values for the excretion of 17-ketosteroids and 17-ketogenic steroids in South African Bantus, and Nigerian males. Simpson (1965) reported differences in urinary 17-hydroxycorticosteroid excretion between European and Equatorial Amerindians. However, urinary steroid levels give an imprecise indication of adrenocortical activity, and the finding that plasma cortisol levels in East African males are similar to those found in other communities (Leonard, 1965) suggests that interracial differences are not important as far as the control of plasma cortisol levels is concerned.

In the present investigation the importance of genetic influences on the pituitary-adrenal axis has been studied by measuring plasma corticosteroid levels in healthy human twin pairs (Becker, 1967).

Material and Methods

Twins Studied. In all, 142 twin pairs from the Glasgow and West of Scotland area volunteered in response to appeals through the local press, radio, and television for healthy twin pairs. Each twin was interviewed and examined simultaneously, in either the afternoon or early evening. None of the twins had clinically apparent disease, nor were any known to be on steroid therapy. All the twins had normal blood pressure readings (Downie et al., 1969). Table I shows the segregation of the 142 twin pairs according to their sex and zygosity. It can be seen that there was a relative excess of monozygotic (MZ) female twin pairs, and that the majority of twins were young. The mean age and age ranges were, however, similar in all groups.

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Determination of Zygosity. The methods used to determine the zygosity of the twins have been described elsewhere (Greig et al., 1967). These methods are not completely accurate but give a likelihood of a correct diagnosis of monozygosity of greater than 90%, in every twin pair accepted as being monozygous in the present study. In the case of dizygous (DZ) twins, however, the likelihood of a correct diagnosis of zygosity was 100%, as zygosity was only diagnosed when there was unequivocal evidence of genetically determined phenotype difference such as variation in facial appearance, fingerprint pattern, hair, or iris colour, or blood group antigen.

Measurement of Plasma Corticosteroids. After determination of blood pressure, venous blood samples were taken from each twin at the same time. The blood was taken into lithium heparin tubes, centrifuged, and the supernatant plasma retained and stored at 0°C. Plasma corticosteroids were measured as the 11-hydroxycorticosteroids using the fluorimetric method described by Mattingly (1962). Though, in man, cortisol is the principal free 11-OHCS in plasma, corticosterone is present in much smaller amounts and is also measured by this method. In this study the term plasma corticosteroids thus refers to the total 11-hydroxycorticosteroids.

Statistical Analysis. The differences between the two members of the pairs of twins are expressed as mean intrapair variances, and can be calculated from the expression \( \sum x^2/n \), where \( x \) is the difference between the two members of a pair of twins (in this instance in the plasma corticosteroid level), and \( n \) is equal to the number of pairs of twins. The method used in this study contrasts the mean intrapair variance of plasma corticosteroid levels observed in MZ (genetically identical) twin pairs, with the mean intrapair variance of plasma corticosteroid levels in the like-sex DZ (genetically non-identical) twin pairs. The variance ratio is calculated from a comparison of the different mean intrapair variances, and its significance is assessed using the F distribution (Osborne and DeGeorge, 1959).

Results

The results are summarized in Tables I, II, and III. The data for unlike-sex dizygotic twin pairs is included in the Tables, but not used in the study, as interpretation of the results is made difficult by the sex difference in this group of twin pairs.

Mean and Range. The mean plasma corticosteroid levels were 14.12 and 14.79 \( \mu g/100 \) ml. for the MZ and DZ female twin pairs, respectively, and 16.22 and 11.96 \( \mu g/100 \) ml. for the MZ and DZ male twin pairs, respectively.

The range of plasma corticosteroid levels in female twins was 5.5-28.5 \( \mu g/100 \) ml. and in male twins 3.5-37 \( \mu g/100 \) ml. (Table I).

Intrapair Variances. The results are first presented for all the male and female twins taken as a whole (Table I), and then analysed separately for male and female twins living together, and those separated for more than one year (Table II).

The intrapair variance for plasma corticosteroids of 12-62 for all the MZ female twin pairs contrasts with the intrapair variance of 32-40 in the DZ female twin pairs. This comparison yields an F ratio of 2.57 which is significant at the less than 1 in a 100 (\( p < 0.01 \)) level (Table III). However, examination of the data for males shows that the intrapair variance for MZ male twin pairs (30.47) was higher than the intrapair variance for the DZ male twin pairs (18.68), which is clearly not significant of a genetic component (Table III).

In Table II the intrapair variance of plasma corticosteroids in twins living together in the same household, and for twins separated from each other for longer than one year, is documented. For

### Table I

<table>
<thead>
<tr>
<th>Twin Group</th>
<th>No. of Twin Pairs</th>
<th>Age (yr.)</th>
<th>Results Plasma Corticosteroid</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
<td>Range</td>
</tr>
<tr>
<td>MZ Female</td>
<td>46</td>
<td>25</td>
<td>10-9</td>
</tr>
<tr>
<td>MZ Male</td>
<td>20</td>
<td>25</td>
<td>10-9</td>
</tr>
<tr>
<td>DZ Female</td>
<td>23</td>
<td>22</td>
<td>15-6</td>
</tr>
<tr>
<td>DZ Male</td>
<td>18</td>
<td>21</td>
<td>10-9</td>
</tr>
<tr>
<td>DZ Unlikely sex</td>
<td>35</td>
<td>19</td>
<td>11-4</td>
</tr>
</tbody>
</table>

* SD, standard deviation.
† See text for explanation of how this function was calculated.
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TABLE II
MEAN INTRAPAIR VARIANCE OF PLASMA CORTICOSTEROID IN TWINS LIVING TOGETHER AND IN TWINS LIVING APART (SEPARATED FOR 1 YEAR OR LONGER)

<table>
<thead>
<tr>
<th>Twin Group</th>
<th>Twins Together</th>
<th></th>
<th>Twins Apart</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No. of Twin Pairs</td>
<td>Age (yr.)</td>
<td>Intrapair Variance† Plasma Corticosteroid</td>
</tr>
<tr>
<td></td>
<td>Mean</td>
<td>Range</td>
<td>SD*</td>
</tr>
<tr>
<td>Female MZ</td>
<td>37</td>
<td>18</td>
<td>17</td>
</tr>
<tr>
<td>Female DZ</td>
<td>5</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>Male MZ</td>
<td>14</td>
<td>16</td>
<td>15</td>
</tr>
<tr>
<td>Male DZ</td>
<td>6</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>DZ Unlike sex</td>
<td>31</td>
<td>16</td>
<td>7-9</td>
</tr>
</tbody>
</table>

* SD, standard deviation.
† See text for explanation of how this function was calculated.

TABLE III
SIGNIFICANCE VALUES FOR OBSERVED INTRAPAIR VARIANCES OF PLASMA CORTICOSTEROIDS IN MALE AND FEMALE TWINS

<table>
<thead>
<tr>
<th>Twin Group</th>
<th>Intrapair Variance*</th>
<th>F Ratio</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>All Twins</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female MZ</td>
<td>12-62</td>
<td>32-40</td>
<td>2-57</td>
</tr>
<tr>
<td>Female DZ</td>
<td>30-47</td>
<td>18-68</td>
<td>0-61</td>
</tr>
<tr>
<td>Male MZ</td>
<td>13-32</td>
<td>40-22</td>
<td>3-05</td>
</tr>
<tr>
<td>Male DZ</td>
<td>27-29</td>
<td>12-89</td>
<td>0-47</td>
</tr>
<tr>
<td>Twins living together</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female MZ</td>
<td>9-72</td>
<td>4-26</td>
<td>0-43</td>
</tr>
<tr>
<td>Female DZ</td>
<td>37-89</td>
<td>33-72</td>
<td>0-88</td>
</tr>
<tr>
<td>Male MZ</td>
<td>30-47</td>
<td>18-68</td>
<td>0-61</td>
</tr>
<tr>
<td>Male DZ</td>
<td>13-32</td>
<td>40-22</td>
<td>3-05</td>
</tr>
<tr>
<td>Twins living apart</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* See text for explanation of how this function was calculated.

The results show a significantly smaller intrapair variance in plasma corticosteroids in MZ as compared to DZ female twin pairs. Before accepting this difference as evidence of genetic control, it should be considered whether the results could be explained by the MZ twins sharing a more uniform environment than did the DZ twins. We consider this alternative explanation to be unlikely in the present study, as the significantly lower intrapair variance for plasma corticosteroid values in MZ females, compared with DZ females, was not matched by the findings in the MZ and DZ male twins, and the vast majority of the twins of both sexes were young and living in the same home environment at the time of study.

It seems reasonable to conclude therefore that the smaller intrapair variance in the female MZ compared to DZ twins almost certainly reflects a fairly strong genetic control of the variation in plasma corticosteroid levels in females. The failure to show a genetic control of plasma corticosteroid levels in males does not mean that such control does not exist, but that it is likely to be weak.

female twin pairs living together the mean intrapair variance for MZ twins was 13-32 and for DZ twins was 40-22. This difference yields an F ratio of 3-05 which is highly significant (p < 0-01) (Table III). When MZ and DZ female twins living apart are compared it is seen that the intrapair variance for MZ females (9-72) is higher than the intrapair variance for DZ females (4-26). There is thus no evidence for genetic control of plasma cortisol in this group, though the numbers are small.

In males living together the intrapair variance of MZ twins was 27-29, while for DZ twins it was only 12-89. There is thus clearly no evidence for a genetic factor. Comparison of intrapair variances in MZ and DZ male twins living apart also failed to show any significant genetic component in the control of plasma corticosteroids. The number of separated and non-separated twins studied was small, but the results were no different from those obtained when all males were studied together.

Discussion

Both genetic and environmental factors have been shown by earlier twin studies to be important in the control of biochemical parameters such as serum cholesterol (Pikkarainen, Takkunen, and Kulonen, 1966) and uric acid (Boyle et al., 1967). To our knowledge, however, there has been no previous study of plasma corticosteroid levels in normal twins.

Our results show a significantly smaller intrapair variance in plasma corticosteroids in MZ as compared to DZ female twin pairs. Before accepting this difference as evidence of genetic control, it should be considered whether the results could be explained by the MZ twins sharing a more uniform environment than did the DZ twins. We consider this alternative explanation to be unlikely in the present study, as the significantly lower intrapair variance for plasma corticosteroid values in MZ females, compared with DZ females, was not matched by the findings in the MZ and DZ male twins, and the vast majority of the twins of both sexes were young and living in the same home environment at the time of study.

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Summary

Plasma corticosteroid levels were estimated by the fluorimetric method of Mattingley in 142 pairs of healthy twins from the Glasgow and West of Scotland area.

Calculation of the intrapair variances in monozygotic and dizygotic twins showed a significant genetic component in the control of variation in the plasma corticosteroid level in females. The study failed to detect a genetic component in males. The suggestion that environmental factors play a more important role in the regulation of plasma corticosteroid levels in males compared with females is discussed.

We are grateful for the co-operation of the many twins who took part in this survey.

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REFERENCES


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