Cytogenetic Studies in a Family with Waldenström’s Macroglobulinaemia

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Waldenström’s macroglobulinaemia is a disease characterized by an increase in the serum of gamma-globulins of high molecular weight (macroglobulin). The chief feature of the bone-marrow of these patients is an infiltration of numbers of small lymphocytes. The clinical and pathological features of primary macroglobulinaemia have been described by several authors since the syndrome was first defined by Waldenström (Waldenström, 1944, 1948; Imhof, Baars, and Verloop, 1959; Mackay, 1959; Martin, 1960; Kok, Whitmore, and Ainsworth, 1963).

A serological and cytogentic study has been carried out on the relatives of a patient with Waldenström’s macroglobulinaemia. The detailed clinical and serological aspects of the family have been presented elsewhere (Brown et al., 1967), and therefore only brief details of these aspects of the family will be given here. The purpose of this article is to describe in full and to discuss the cytogentic abnormalities found in this family, particularly with regard to the incidence of pseudodiploidy.

Material and Methods

Paper electrophoresis of the serum proteins was carried out by the method of Tiselius (1939) and immunoelectrophoresis by the method of Scheidegger (1955), using horse anti-human serum (Pasteur Institute). Antiglobulin neutralization tests (Kekwick et al., 1961) were carried out, using a reagent containing anti-γ1 and anti-γ2, and red cells sensitized with anti-D to detect γ2-globulin, and cells coated with anti-Lea in the absence of complement (Stratton, Gunson, and Rawlinson, 1962). Sia test was carried out by the method of Martin (1960). Blood samples were taken from available members of the family and chromosomes were obtained from cultured lymphocytes, according to the method of Hungerford et al. (1959). All cells containing the abnormal marker chromosome were examined directly with the microscope and then from

II. 1, aged 62 years. This female patient showed the characteristic features of Waldenström’s macroglobulinaemia. The onset of the illness was insidious, with weight loss, lassitude, anaemia, spontaneous bleeding, and cardiac failure. The diagnosis was suggested by a greatly increased erythrocyte sedimentation rate and by the presence of a dense band in the γ-region on paper electrophoretic examination of the serum. The Sia water test was positive. Immunological studies and ultracentrifugal examination confirmed the diagnosis.

The patient died three years after the initial examination and necropsy showed the typical, diffuse, pleomorphic infiltrate of the marrow and lymph nodes.

Six relatives were screened by clinical examination, routine laboratory tests, and serological studies, and one relative (II. 2) showed a serum protein abnormality. A second relative (III. 5) showed a slight increase in γ-globulin. Details are shown in Table I.

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### TABLE I

<table>
<thead>
<tr>
<th>Case</th>
<th>Hb (g./100 ml.)</th>
<th>WBC per c.mm.</th>
<th>ESR mm./hr. (Westergren)</th>
<th>Latex Agg’n Tests</th>
<th>Blood Group</th>
<th>Total Serum Proteins (g./100 ml.)</th>
<th>Serum Alb. (g./100 ml.)</th>
<th>Serum Glob. (g./100 ml.)</th>
<th>Sia Test</th>
<th>Paper Electrophoresis</th>
<th>W.R.</th>
<th>Immuno-electrophoresis</th>
<th>Antiglob. Neutral’n Test</th>
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</thead>
<tbody>
<tr>
<td>II. 2</td>
<td>12-6</td>
<td>4400</td>
<td>38</td>
<td>+ve +ve</td>
<td>A Rh +ve</td>
<td>7-7</td>
<td>3-7</td>
<td>4-0</td>
<td>-ve</td>
<td>Inc’d band γ-region</td>
<td>Normal</td>
<td>Normal</td>
<td>γγ-glob. inc’d</td>
</tr>
<tr>
<td>III. 2</td>
<td>13-4</td>
<td>3200</td>
<td>3</td>
<td>-ve -ve</td>
<td>A Rh +ve</td>
<td>6-9</td>
<td>3-9</td>
<td>3-0</td>
<td>-ve</td>
<td>Normal</td>
<td>γγ-glob. band</td>
<td>Normal</td>
<td>Normal</td>
</tr>
<tr>
<td>III. 2</td>
<td>13-0</td>
<td>4200</td>
<td>2</td>
<td>-ve -ve</td>
<td>A Rh +ve</td>
<td>6-8</td>
<td>4-0</td>
<td>2-8</td>
<td>-ve</td>
<td>Normal</td>
<td>Normal</td>
<td>Normal</td>
<td>Normal</td>
</tr>
<tr>
<td>III. 1</td>
<td>13-4</td>
<td>6800</td>
<td>1</td>
<td>-ve -ve</td>
<td>A Rh +ve</td>
<td>6-5</td>
<td>3-8</td>
<td>2-7</td>
<td>-ve</td>
<td>Normal</td>
<td>Normal</td>
<td>Normal</td>
<td>Normal</td>
</tr>
<tr>
<td>III. 4</td>
<td>12-7</td>
<td>5400</td>
<td>2</td>
<td>-ve -ve</td>
<td>O Rh +ve</td>
<td>6-2</td>
<td>3-6</td>
<td>2-6</td>
<td>-ve</td>
<td>Normal</td>
<td>Normal</td>
<td>Normal</td>
<td>Normal</td>
</tr>
<tr>
<td>III. 5</td>
<td>11-7</td>
<td>6200</td>
<td>3</td>
<td>-ve -ve</td>
<td>A Rh +ve</td>
<td>6-8</td>
<td>3-6</td>
<td>3-2</td>
<td>-ve</td>
<td>Slightly inc’d band γ-region</td>
<td>Normal</td>
<td>Normal</td>
<td>Normal</td>
</tr>
</tbody>
</table>

#### II. 2, aged 58 years.

This female patient had a history of encephalitis two years before the present study. At that time serum protein estimation was normal and the erythrocyte sedimentation rate was 28 mm. in one hour (Westergren). Two years later the total serum proteins were 7.7 g./100 ml. with a serum globulin level of 4 g./100 ml. The erythrocyte sedimentation rate was 38 mm. in one hour (Westergren), and paper electrophoresis of the serum showed a dense band in the γ-region. Immuno-electrophoretic and antiglobulin neutralization tests (kindly performed by Dr. H. H. Gunson of the Blood Transfusion Service, Manchester) showed an increase in γγ-globulin.

No clinical or serological abnormalities were found in the remaining five relatives who were investigated.

#### Cytogenetic Data

II. 1 had 2 cells with 47 chromosomes, the additional autosome being larger than any others in the karyotype and typical of that found in some other cases of macroglobulinaemia. In one cell this abnormal chromosome was metacentric and in the second it was submetacentric (Fig. 2). Two pseudodiploid cells were also found, which contained a large submetacentric chromosome. In one of these cells a No. 21–22 chromosome was missing, while in the second a No. 2 chromosome was absent. In this latter cell it was thought that the macroglobulinaemia chromosome (M chromosome) was the missing No. 2, but on comparing them, the M chromosome was 26% longer than the No. 2. The incidence of the abnormality in this patient was therefore 8%.

In II. 2 no cells containing a typical macroglobulinaemia chromosome complement were seen (i.e. 47 chromosomes, the supernumerary autosome being either a large metacentric or submetacentric chromosome). There were however a number of pseudodiploid cells which contained a large submetacentric chromosome. In two of these a No. 2 chromosome was absent and the M chromosome was 22% larger than the No. 2 present; they were therefore not considered to be homologous, as they were also morphologically dissimilar. Another cell had a submetacentric M chromosome, but one of the small autosomes of group 19–20 was absent. In a fourth cell a similar M chromosome was present, but a small member of group 6–12 (M 12) was absent. 16% of the cells, therefore, carried an M chromosome.

In III. 3, no abnormal cells were seen.

In III. 1, no typical 'macroglobulinaemia' karyotype was found. One pseudodiploid cell had a submetacentric M chromosome, but a No. 2 autosome was absent. Again the M chromosome was not considered to be the missing No. 2, as it was 28% longer. Two cells had a similar M chromosome, but a small member of group 6–12 (M 12) was missing. 3% of the cells were, therefore, abnormal.

In III. 2, two cells with 47 chromosomes containing an additional large chromosome were seen. In one cell

![Fig. 2. Chromosomes of groups 1–5 from a cell containing a submetacentric M chromosome.](http://jmg.bmj.com/ on August 14, 2017 - Published by group.bmj.com)
RESULTS OF CHROMOSOME COUNTS IN FAMILY

<table>
<thead>
<tr>
<th>Case</th>
<th>Distribution (No. of mitoses)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>44</td>
</tr>
<tr>
<td>III. 1</td>
<td>3</td>
</tr>
<tr>
<td>II. 2</td>
<td>1</td>
</tr>
<tr>
<td>III. 1</td>
<td>2</td>
</tr>
<tr>
<td>III. 2</td>
<td>1</td>
</tr>
<tr>
<td>III. 3</td>
<td>1</td>
</tr>
<tr>
<td>III. 4</td>
<td>1</td>
</tr>
<tr>
<td>III. 5</td>
<td>2</td>
</tr>
</tbody>
</table>

TABLE II
RESULTS OF CHROMOSOME COUNTS IN FAMILY

Discussion

Apart from the propositus only one other member of the family has an abnormal serum globulin pattern, in which the $\gamma_1$ fraction was raised. III. 5 had a slight increase in gammaglobulin on electrophoresis, but this has not been confirmed by the antiglobulin neutralization test. We cannot therefore at present regard him as abnormal.

There are two studies of familial serum protein abnormalities associated with macroglobulinaemia (Massari, Fine, and Metais, 1962; Seligmann and Badin, 1962). No cytogenetic studies have been made in such families as far as we are aware. Chromosome abnormalities have now been found in a number of cases of Waldenström's macroglobulinaemia (Bottura, Ferrari, and Veiga, 1961;
German, Biro, and Bearn, 1961; Benirschke, Brownhill, and Ebaugh, 1962; Elves and Israëls, 1963). There is, however, no specific abnormality. The abnormal chromosome which is present in only a small proportion of dividing lymphocytes may be either a large metacentric, or a submetacentric chromosome. In one case only is the additional large chromosome almost acrocentric (Bottura et al., 1961). Similar abnormalities may be found occasionally in subjects with mild degrees of hypergammaglobulinaemia without any evidence of macroglobulinaemia (Elves and Israëls, 1963).

The abnormal chromosome may therefore exist without any abnormal protein fraction being produced. This family is another example of this phenomenon. It is also the first in which the chromosomal abnormality has been shown to be familial. The abnormal chromosome is present in 5 out of the 7 subjects studied, and in all but one cell it is the large submetacentric variety (Type II). One metacentric chromosome (Type I) has been found in a cell from the propositus. Only in the propositus is there any evidence of excess macroglobulin production.

Of interest in this family is the incidence of pseudodiploidy. In a number of cells the abnormal chromosome is present, but the chromosome number remains modal (i.e. 46) due to the loss of one of the normal autosomes. It is questionable how much attention should be paid to these cells, particularly when the missing autosome is one of the smaller members of the karyotype. The absence of other similar cells, either from the same subject or from others, suggests that these cells may have suffered loss of a chromosome during the preparation of the chromosomes for study. Thus, it is perhaps more desirable to consider these cells as 'normal' macroglobulinaemia cells and not additional abnormal stem cell lines. Some pseudodiploid cell lines do appear consistently, however, both in different cells of the same subject and also in cells of different subjects. One of these abnormal cell lines possesses the Type II chromosome, but is deficient in a normal No. 2 autosome. The M chromosome was considered to be the second No. 2 autosome until it was found that it was consistently longer, the difference in size ranging from 16–28% of the mean length of the pair. As controls, cells from 10 normal unrelated subjects were examined and the percentage difference in length of the No. 2 homologues was estimated. The range of difference found was 0–11%, with a mean of 4%. This type of cell has been found in all of the abnormal subjects, including the propositus (Table II). The second abnormal pseudodiploid cell type is deficient in a smaller member of group 6–12. This cell type is present in 4 subjects. We are reluctant to dismiss these cells as artefacts. Furthermore, Houston, Ritzmann, and Levin (1967), studying a series of patients with macroglobulinaemia, have found a high incidence of structural or numerical abnormalities in one of the smaller pairs of chromosomes in group 6–12. They also suggest that the No. 12 pair was involved in these abnormal cells.

As the abnormal chromosome in these blood cultures is confined to only a small proportion of cells of the lymphoid series, it is clear that the abnormal stem cell line originates during the later period of embryonic growth or in postnatal development. It is as yet uncertain whether or not it is only the lymphoid cell types that carry this abnormality, or whether there is a general mosaicism throughout all cells of the body. So far no examination has been made of the karyotype of skin cells in such patients, and we were unable to carry out this investigation in this family. In addition, it was not possible to obtain bone-marrow specimens from the relatives of the propositus. There is some evidence, however, that abnormal stem cells may be present in the bone-marrow. Bottura et al. (1961) reported the presence of a large chromosome in some of the cells obtained from a patient with this disease by sternal puncture, and others have also found abnormal cells in the marrow of patients with macroglobulinaemia (Heni and Siebner, 1963; Houston et al., 1967).

The bone-marrow in this condition, however, usually contains increased numbers of small lymphoid cells, and it may be these cells or their precursors that carry the abnormal chromosome rather than cells of the erythrocyte or granulocyte series. Of interest in this context is the patient described by Houston et al. (1967) who, in addition to macroglobulinaemia, also had chronic myeloid leukaemia. Cells containing the abnormal chromosome were found in both blood cultures and direct bone-marrow samples from this patient. The authors were not able to identify with certainty the cell type represented by the abnormal mitotic figures, but the majority of cells in the marrow were of the myeloid series. Further, Ph1 containing cells, which are thought at present to be non-lymphocytic, were also seen and contained the abnormal 'macroglobulinaemia' chromosome. Thus, it appears that the abnormal cell type is not necessarily confined to cells of the lymphoid series. It may be suggested that the pathological serum protein pattern in these patients is due to the existence in the lymphoid population of a 'clone' of cells which contain excess
genetic material controlling the synthesis of gammaglobulin (carried by the abnormal chromosome). Its presence in three members of this family in which no protein abnormality was detectable indicates that gammaglobulin overproduction is not an inevitable consequence of the presence of this chromosome. It can be postulated that the abnormal cell line may lie dormant until a suitable stimulus is encountered which normally induces gammaglobulin production, and, concomitant with this, abnormal gammaglobulin would also be formed. The nature of such a stimulus is not clear, but as there is evidence that the lymphocyte is concerned in immunological processes, it may be antigenic in nature. The encephalitis suffered by II.2 may have provided such a stimulus. At the time of her illness, this patient had normal gammaglobulin levels which are now raised.

An alternative hypothesis is that the abnormal cell line is suppressed by a homeostatic mechanism. If there was a breakdown of this immuno-suppressive mechanism, the abnormal cells would be able to proliferate. Phytohaemagglutinin, if it acted in a manner suggested by Coulson and Chalmers (1964), would short-circuit the recognition mechanism and would stimulate all immunologically competent cells to transform and divide. Thus, the blood culture technique provides an artificial system in which cell lines not normally active may be revealed.

Summary

Chromosome abnormalities were demonstrated in 4 out of 6 relatives of a patient with Waldenström's macroglobulinaemia. Apart from the proposition, only one other member of the family had a significantly abnormal serum protein abnormality in which the $\gamma_1$ fraction was increased. The abnormal cells contained a large submetacentric chromosome, except for one large metacentric chromosome which was found in a cell from the proposition.

Encephalitis was followed by a gammaglobulin abnormality in one of the relatives. An antigenic stimulus may initiate gammaglobulin overproduction or, alternatively, proliferation of abnormal cells may result from a disturbance of the normal immuno-suppressive mechanism.

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