Multiple myeloma in Jamaica: A study of 40 cases with special reference to the incidence and laboratory diagnosis

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SYNOPSIS For two years protein abnormality was studied in 40 cases of myelomatosis in Jamaica. Thirty-nine of these were in West Indian Negroes. The minimum incidence of myelomatosis in this group was estimated to be of the order of 50 cases per million per annum which is considerably higher than in Caucasians as reported by previous workers. A larger number showed myeloma protein with β-globulin mobility and hypogammaglobulinaemia than with γ-globulin mobility. As in the Caucasians the disease is more common in men than in women and the age incidence in both seems to be the same. Combined serum and urinary electrophoresis was diagnostic in every case, and examination of the urine for Bence-Jones protein by electrophoresis yielded more consistent findings than the classical heat test. It is suggested that combined serum and urine electrophoresis should be done in all cases of suspected myelomatosis.

Since electrophoresis was first used as an aid in the diagnosis of multiple myeloma in 1939 by Longsworth, Shedlowsky, and MacInnes, several reviews and several series on myelomatosis in Caucasians have been published. However, very little is known about the incidence of myeloma in the Negro race. Isolated cases have been published, and in a survey of myelomatosis in 34 Brooklyn hospitals, New York, MacMahon and Clark (1956) noted a higher incidence in Negroes than in Caucasians.

The occurrence of hypoproteinaemia and hypogammaglobulinaemia in myelomatosis is unusual and sometimes contributes to the difficulty experienced in the differential diagnosis of multiple myeloma, especially if either the bone marrow biopsy or the radiological appearance is inconclusive.

In the present series an attempt is made to study the incidence of multiple myeloma in the Negro race in Jamaica and to point out some important aspects concerning the laboratory diagnosis of myelomatosis, especially where the serum protein electrophoresis appears normal or shows only a hypogammaglobulinaemia.

MATERIALS AND METHODS

The material consisted of 40 cases in which myelomatosis was well documented. With the exception of one, all were Negroes. These were investigated in the course of our routine work in the Sub-department of Chemical Pathology in the University of the West Indies between 1962 and 1964. Twenty-five of these were men whose average age was 54 years (range 40-71 years). The 15 women had an average age of 50 years (range 40-65 years). The average for the whole group was 52 years, and Table I shows the age incidence of myelomatosis in Jamaica.

<table>
<thead>
<tr>
<th>Age (yr.)</th>
<th>No. of Males</th>
<th>No. of Females</th>
</tr>
</thead>
<tbody>
<tr>
<td>31-40</td>
<td>5</td>
<td>3</td>
</tr>
<tr>
<td>41-50</td>
<td>9</td>
<td>6</td>
</tr>
<tr>
<td>51-60</td>
<td>6</td>
<td>5</td>
</tr>
<tr>
<td>61-70</td>
<td>3</td>
<td>1</td>
</tr>
<tr>
<td>71-80</td>
<td>2</td>
<td>Nil</td>
</tr>
<tr>
<td>Total</td>
<td>25</td>
<td>15</td>
</tr>
</tbody>
</table>

Electrophoresis by the method described by Varley (1962) was carried out on filter paper and also on cellulose acetate according to the method of McFarlane (1963). Both methods gave identical results. Samples of urine, each of 5 ml., were concentrated for approximately 90 minutes and electrophoresed according to the method of McFarlane (1964). Urine electrophoresis for Bence-Jones protein instead of the classical heat test was adopted since it proved more reliable than the latter. Total protein and globulin were estimated both by the
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RESULTS

TOTAL PROTEIN As shown in Fig. 1, of the 36 patients whose total protein was estimated, four (11% of the total) had hypoproteinaemia with values between 4·0 and 6·0 g./100 ml; 17 patients (47% of the total) had normal total protein concentration, that is, of values between 6·1 and 8·0 g./100 ml, whereas 15, or 42%, had hyperproteinaemia with values between 8·1 and 14·0 g./100 ml.

TOTAL GLOBULIN Figure 1 also shows the total globulin levels of the patients studied. Here it can be seen that nine cases, 25% of the patients, had relative hypoglobulinaemia, with values between 1·0 and 3·0 g./100 ml; four cases (11%) had normal globulin values between 3·1 and 4·0 g./100 ml., and 23 cases (64%) had hyperglobulinaemia with values ranging from 4·1 to 11·0 g./100 ml.

SERUM ELECTROPHORESIS This was performed on all the sera. Sixteen cases showed a sharp beta globulin peak with hypogammaglobulinaemia, 14 a typical gamma globulin peak, in four cases there was a definite peak in the alpha region, and six cases gave a normal serum protein pattern on electrophoresis. When the urines of the latter were concentrated and electrophoresed a dominant band of globulin was invariably found in the \( \beta - \gamma \) region, obviously the Bence-Jones protein.

URINE ELECTROPHORESIS This was performed on 16 cases and in 13 of these the electropherogram of the concentrated urine was diagnostically important, especially (a) in those cases where the
serum electrophoresis was normal or nearly so; 
(b) in those with hypogammaglobulinaemia; and 
(c) in those patients whose diagnosis was missed by 
the clinicians. The urine electrophoretic pattern of 
the six cases which had a normal serum electrophoretic pattern showed that all the myeloma 
protein seemed to be excreted in the urine, some-
times accompanied by a small amount of both 
albumin and other globulins. Figure 2 shows typical 
electropherograms of both urine (U) and serum 
(S) from three patients with multiple myeloma. The 
serum electropherograms of both A and B show 
only minor non-specific changes, slight hypogamma-
globulinaemia, and no detectable myeloma proteins, 
whereas the urine electropherograms in these cases 
show that all the abnormal protein is found in the 
urine. Although the Bence-Jones protein in A is 
accompanied by some albuminuria, the diagnosis is 
clear, since the dominant globulin band is obvi-
ously a Bence-Jones protein. Similarly in the urine 
electropherograms shown in Fig. 2, B and C show a 
single dominant band in the beta globulin region 
which is clearly a Bence-Jones protein. In Fig. 2C 
the Bence-Jones protein can be seen migrating in the 
same slow gamma globulin region as the serum 
myeloma protein.

DISCUSSION

Although specific comparative data are few, the 
finding of 39 cases of myelomatosis in Negroes in 
Jamaica within two years seems high. This figure 
is of the order of 20 cases per annum, and as it has 
since been estimated that our routine laboratory 
serves a population of not more than 400,000 people, 
this is equivalent to 50 cases per million per annum. 
The figure of 400,000 is an overestimation, since there 
are more than two other routine laboratories of 
equivalent size serving the same population. 
Furthermore there must be several cases of myeloma 
that go undiagnosed each year, due to the shortage 
of medical facilities in the Island. Clearly, then, the 
incidence of myeloma appears to be higher than the 
figure indicated here. MacMahon and Clark (1956), 
on the basis of a survey of 279 cases of myeloma in 
34 Brooklyn hospitals in New York in the period 
1943-53, reported the incidence of multiple myeloma 
in Negroes as 23·5 per million per annum which is 
more than twice the rate in whites in the same 
area, and the excess risk experienced by the non-
white compared with the white population is 
probably closer to 100% than 20%. Comparative 
statistics on the incidence of multiple myeloma in 
other countries are rare. However, Heinivaa and 
Eisalo (1960) saw 34 cases of myeloma from 1954 to 
1959 at the Second Department of Medicine, 
University of Helsinki. Ramot and Salomi (1961) 
reported that the frequency per 100,000 of myeloma 
for the years 1956-60 was found to be 4·69 for the 
European-American born Jews and 3·65 for the 
found 61 cases of multiple myeloma at Malmö, 
Sweden, from a population of 200,000 during the 
10 years between 1950 and 1959. This is equivalent 
to about 30 cases per million per annum, which is 
significantly lower than the annual figure of the 
present myeloma series in Negroes in Jamaica and 
is of the same order as the figure of 26 cases per 
million per annum for England (Martin, 1961). 
The significance of the high incidence of mye-
ломatosis in Negroes is not known, although it is 
tempting to suggest that some genetic factor may 
be involved.

In the present series 62% of the total cases were 
males. This is in agreement with the findings of 
other workers (Snapper, Turner, and Moscovitz, 
1953; Martin, 1961). All that can be said concerning 
the mean age incidence of 52 years for the present 
series is that it appears to be similar to that reported 
for Caucasians by previous workers. 

A significant proportion of the cases, 25%, in the 
present myeloma series, had a low total protein and 
globulin level. On electrophoresis it was found that 
as many as 60% (15% of the total number of cases) 
of these gave a normal pattern or a pattern with only 
minor non-specific serum electrophoretic changes. 
On account of the normal serum electrophoretic 
pattern in these cases the laboratory diagnosis of 
multiple myeloma would most certainly have been 
missed if the urines had not been electrophoresed 
simultaneously.

In the present series the occurrence of the myeloma 
protein in the β-globulin region together with 
hypogammaglobulinaemia was more common than in 
the γ-globulin region, 40% of the former to 30% 
of the latter. In the literature the highest percentage 
of myelomas is usually found with slow gamma 
globulin mobility but the significance is not known. 
Alpha myelomas are extremely rare, and the 
presence of four in this series is indeed interesting. In 
all four cases there was a single sharp peak in the serum 
α2 region. Two of these patients excreted Bence-
Jones protein and in one of these the Bence-Jones 
protein had the same mobility as the corresponding 
serum myeloma protein.

However, Waldenström (1952), Osseman (1959), 
and others advise that myelomas should only be 
classified as α2 with great caution, since it is well 
known that generalized carcinomas and infections 
which are so common in multiple myeloma may 
give rise to a marked increase of alpha globulins. 
Whenever a low globulin level with a normal
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serum electrophoretic pattern was encountered in a suspected case of multiple myeloma, attempts were specially made to obtain a specimen of urine from such patients for electrophoretic examination. Invariably, simultaneous electrophoresis of such sera and urines of these patients proved to be diagnostic even when the heat test for Bence-Jones protein was negative. During this study it was found that simultaneous electrophoresis of the serum and urine was the first means of detecting multiple myeloma in no less than 20% of the cases. These were subsequently confirmed either by bone marrow biopsy or by x-ray studies or both. Recently Clough and Reah (1964) observed that over a three-year period 15 cases of multiple myeloma had been detected in the Biochemistry Department entirely by careful examination of urinary and serum proteins.

More conclusive diagnostic results were obtained when urines from suspected patients were rapidly concentrated and electrophoresed on cellulose acetate than from attempting to do the heat test for Bence-Jones protein. Osserman and Lawlor (1955) reported three cases which gave negative results by the heat test but a positive reaction for Bence-Jones protein was found on urine electrophoresis. Owen and Rider (1957) showed that the two main criteria on which the usual heat test for Bence-Jones protein depend are unreliable, and Burtin (1964) stated that urinary electrophoresis provides a more reliable means of evaluation, when it discloses the existence of a unique urinary protein with \( \beta-\gamma \) or intermediate mobility.

The results presented here confirm the findings of previous publications (Owen and Rider, 1957; Osserman, 1959; Waldenström, 1962) that about 85% of myeloma sera show the typical 'church spire' pattern on electrophoresis; the other 15% may show only hypogammaglobulinaemia or a normal pattern or an electrophoretic pattern with minor non-specific changes. However, in this 15% of patients when the urine is electrophoresed simultaneously with the serum diagnostic results have been obtained. It is suggested, therefore, that both serum and urinary electrophoresis should be performed in all cases of suspected myelomatosis, particularly those with abnormal or low total serum protein. No false positive results were obtained by this procedure, and in fact it should be pointed out that one was able to rule out a diagnosis of multiple myeloma in several patients when their combined serum and urine showed no characteristic pattern. It was also found that this procedure was especially useful in those cases of multiple myeloma in which inconclusive findings were obtained in either the radiological or bone marrow biopsy investigations or both.

I wish to thank Professor Bras, Head of the Department of Pathology, University of the West Indies, for his interest in this work; Mrs. Jackie Chang, and other technicians of the Sub-department of Chemical Pathology for technical assistance, Dr. L. M. Burke, St. James Hospital, Montego Bay, for his assistance in sending me information on some of the patients, and Mr. D. D. Simmonds, University of Ibadan, for his assistance with the diagrams.

REFERENCES


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doi: 10.1136/jcp.19.3.268