Immunoglobulin-bearing polymorphonuclear leucocytes in primary syphilis

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SUMMARY Non-specific defence mechanisms are thought to be ineffective in eliminating Treponema pallidum after infection in the early stages, but studies aimed at the elucidation of this phenomenon are few.

In this study, the number of IgG-bearing polymorphonuclear leucocytes in patients with primary syphilis was increased; this is most probably a normal phenomenon of infection.

Introduction

The non-specific defence mechanisms are generally thought to be ineffective in eliminating Treponema pallidum after infection has been established. The combined efforts of polymorphonuclear leucocytes, mononuclear macrophages, and non-specific humoral factors are unable to neutralise the invading micro-organisms in the early stages of syphilis. Studies of polymorphonuclear leucocytes (PMNL) in syphilis are few. Histologically, PMNL are seen in the initial stage of primary syphilis, accumulating in the perilymphatic spaces, since later they are replaced by lymphocytes and plasma cells.1

Recently the elimination of treponemes by phagocytes after 7-8 hours of penicillin therapy could be demonstrated in early syphilitic lesions in humans using the electron microscope.2 In another study, syphilitic infection of rabbits was found to be significantly associated with increased metabolic activity of phagocytes as detected with the nitroblue tetrazolium test.3

In this study, the number of immunoglobulin-bearing PMNL, isolated from peripheral blood, in a group of patients with primary syphilis was determined. Variations in the percentages of different Ig classes present on the membrane of circulating PMNL have previously been found in diseases such as allergic contact dermatitis and psoriasis.4

Patients and methods

STUDY POPULATION

The patients in this study were referred by the outpatient department of our hospital. A total of seven patients, all with primary syphilis, were studied. This group of patients was the same as that described in two previous papers on the percentages of B and T lymphoid cells in these patients.5 6 All the patients were male; five were untreated at the time of investigation whereas two were treated for less than one week (patients 9 and 12). The time interval from date of infection to time of investigation varied from 4-12 weeks.

HEALTHY CONTROLS

Ten healthy staff members served as controls for the quantitative determination of immunoglobulin-bearing PMNL. All were male and of the same age group as the patients studied.

ISOLATION OF POLYMORPHONUCLEAR LEUCOCYTES

Separation of PMNL was carried out according to the method of Böyum7 with some minor modifications as described below. Fifty millilitres of peripheral blood were collected aseptically in Erlenmayer bats and defibrinated by shaking for 10 minutes with 50 glass beads; to this, 20 ml of phosphate-buffered saline (PBS) was added. This suspension was layered on top of 40 ml Isopaque-Dextrans* for 45 minutes; the red cells are clumped at

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In-Vitro Immunofluorescence Studies of PMNL

After the isolated PMNL had been washed three times in PBS at 200 x g for five minutes, the relative percentages of IgG-, IgA-, IgM-, IgD-, and IgE-bearing PMNL were determined by immunofluorescence as described before.

Peripheral Blood Counts

Total peripheral leucocyte counts and their differentiation were estimated to relate percentage figures to the absolute numbers of PMNL.

Statistical Method

The non-parametric Wilcoxon’s test with the usual correction for ties, and the normal approximation to obtain the P value, was used for statistical analysis of the numbers of immunoglobulin-bearing PMNL in patients with primary syphilis and in healthy controls.

Materials

The materials used in this study were the same as those used in a previous study on lymphoid cells.

Results

The results of the determination of percentages of immunoglobulin-bearing polymorphonuclear leucocytes for each of the five subclasses of immunoglobulins in patients and controls are given in the table. Absolute numbers of PMNL in the patients with primary syphilis were normal (mean 4.41 x 10^9/l; range 1.39-8.38 x 10^9/l) and statistical analysis was performed on percentage figures. The exact values of standardised W and two-sided tail probability, resulting from applying Wilcoxon's test to the obtained percentages, are also given in the table.

Although the number of patients studied was small, the percentage of IgG-bearing PMNL was significantly increased in the patients with primary syphilis (P = 0.0032). Furthermore, the number of IgE-bearing PMNL were significantly decreased in patients with primary syphilis (P = 0.0206) as were the number of IgD-bearing PMNL (P = 0.0140).

TABLE Percentages of immunoglobulin-bearing polymorphonuclear leucocytes in seven patients and 10 controls

<table>
<thead>
<tr>
<th>Immunoglobulin-bearing PMNL (%)</th>
<th>IgG</th>
<th>IgA</th>
<th>IgM</th>
<th>IgD</th>
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Wilcoxon's test

Probability 0.0032 0.0983 1.1761 -2.0700 -2.3640

Discussion

In this study, data are presented which suggest alterations in surface immunoglobulins of PMNL as a result of infection with T pallidum. The increased number of IgG-bearing PMNL in some but not all patients with primary syphilis may indicate adherence of immune complexes consisting of treponemal antigens and antitreponemal IgG, these complexes being bound by way of C3b and Fc receptors to the membrane of circulating PMNL. Phagocytosis and intracellular processing of treponemal constituents must have taken place since humoral responses were generally present in the patients studied.

Interpretation of the decreased number of IgD- and IgE-bearing PMNL is difficult and its significance too low to allow conclusions to be drawn. Increased numbers of IgG-bearing PMNL do not seem to be disease-specific but seem to be the result of IgG-production as a humoral response to infection. In this study, no data on the ineffectiveness of non-specific defence mechanisms in syphilis are given but the results may indicate the normal processing of immune complexes in primary syphilis. Further studies to elucidate the origin of ineffective non-specific defence mechanisms in syphilis are needed.
References


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