Spontaneous cytotoxicity of rheumatoid and normal peripheral blood mononuclear cells against 4 human lymphoblastoid cell lines

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SUMMARY By measuring spontaneous cytotoxicity of unfractionated peripheral blood mononuclear cells from patients with rheumatoid arthritis and from normal individuals against 4 human lymphoblastoid cell lines we have been unable to demonstrate any preferential recognition of antigens on these cell lines by rheumatoid patients.

Man has a lymphocyte population which is spontaneously cytotoxic to in-vitro propagated tumour cell lines. This phenomenon is mediated by a non-B, non-T, Fc receptor bearing lymphocyte, the spontaneous killer (SK) cell.\(^1\) A similar phenomenon in mice is mediated by the natural killer cell (NK) cell.\(^2\) The biological significance of these systems is unknown, but postulated roles have included tumour surveillance and defence against viral infection. The nature of the target antigen which these cells recognise is unknown. However, SK cells preferentially kill some lymphoblastoid cell lines, especially those derived from tumour cells rather than by viral transformation of normal lymphocytes,\(^3\) and inhibition experiments in mice have implied a certain amount of specificity.\(^4\) Mechanisms controlling SK cells are not known, but Jondal and Targan\(^5\) have postulated that antigens like those expressed on Epstein-Barr virus (EBV) transformed B cell lines may be involved in the in-vivo generation of SK cells.

There is continuing interest in the possibility that rheumatoid arthritis may be caused by a virus. This has been heightened by the observation that about 70% of patients with rheumatoid arthritis have a serum antibody reacting with a nuclear antigen within lymphoblastoid cell lines containing EBV genome.\(^6\) Thus, since it is possible that antigens similar to those expressed on EBV transformed cell lines might be involved in the in-vivo generation of SK cell activity, and since there is evidence of a serological response in rheumatoid arthritis against nuclear antigens in EBV-containing lymphoblastoid cell lines, we decided to investigate SK cell activity against lymphoblastoid cell lines in rheumatoid patients.

**Materials and methods**

**Patients** The results presented are derived from testing peripheral blood mononuclear cells (PBM) from 51 patients with rheumatoid arthritis, 31 females and 20 males, mean age 57.2 years. All were suffering from definite or classical rheumatoid arthritis by ARA criteria\(^7\) except 2 who had probable rheumatoid arthritis. Thirty-four patients were seropositive and 17 seronegative. Most had severe active disease as reflected in a mean ESR of 35.1 and the fact that all except 17 patients were being treated with gold, D-penicillamine, or prednisone. Thirty-seven controls were tested, 20 females and 17 males, mean age 40.5, who were attending for treatment of non-inflammatory rheumatological conditions, mainly back pain.

**Cell lines** Four cell lines were used in these experiments. These were selected to include B and T cell lines, lines positive and negative for EBV genome, and lines derived from tumour cells and by transformation of normal lymphocytes. The main characteristics of these cell lines are shown in Table 1. CCRF/CEM is an EBV genome negative T cell line established from a patient with lymphocytic leukaemia in childhood.\(^8\) RAJI is a well known EBV genome positive B cell line established from Burkitt’s
lymphoma.\textsuperscript{9–11} It is positive for Epstein-Barr virus nuclear antigen (EBNA) but does not express viral capsid antigen (VCA) or membrane antigen (MA) under normal circumstances.\textsuperscript{12} RPMI 8226 is an EBV genome negative B cell line established from a patient with myeloma and produces free immunoglobulin light chains in culture.\textsuperscript{13} Wil is an EBV genome positive B cell line established from normal lymphocytes obtained from a patient with myeloid leukaemia at the Queensland Institute of Medical Research (hence QIMR Wil).\textsuperscript{14}

**CELL CULTURE**

All cell lines were grown in static culture in Eagles suspension medium (Gibco) buffered with Hepes and bicarbonate and supplemented with 10% fetal calf serum, 2 mM L-glutamine and antibiotics (gentamicin 40 \(\mu\)g/ml and amphotericin B 250 \(\mu\)g/ml). All the cell lines except RPMI 8226 grow in suspension and were harvested twice weekly by removal of half their volume. RPMI 8226, which grows as an adherent monolayer, was harvested twice weekly by vigorous shaking and removal of a majority of the culture fluid.

**MEDIUM USED FOR ASSAYS**

Assays were carried out in Eagles MEM (Gibco), buffered with Hepes and bicarbonate and supplemented with 10% fetal calf serum, 2 mM L-glutamine and gentamicin 40 \(\mu\)g/ml (TCM). The same batch of fetal calf serum was used throughout.

**LABELLING OF TARGET CELLS**

After one wash in fresh TCM 5–8 \(\times\) 10\(^6\) lymphoblastoid cells were suspended in 0.1 ml \(^{51}\)Cr-sodium chromate solution (Radiochemicals, Amersham, CJS-IP, specific activity 1 mCi in 1 ml). The mixture was incubated for one hour at 37°C, cells washed 4 times in TCM and adjusted to a concentration of 3 \(\times\) 10\(^8\)/ml.

**SEPARATION OF PERIPHERAL BLOOD MONONUCLEAR CELLS**

Venous blood was collected in sterile containers with 10 U heparin per ml of blood. Peripheral blood mononuclear cells (PBM) were harvested from Triosil/Ficoll gradients.\textsuperscript{15} Monocytes were not routinely removed.

**CYTOTOXIC ASSAY**

Assays were carried out in triplicate in 2.5 ml capped plastic tubes (Seward Co. Ltd.) in a total volume of 400 \(\mu\)l TCM containing 3 \(\times\) 10\(^4\) target cells and 6 \(\times\) 10\(^6\) lymphocytes, a ratio of 20:1 effector to target cells. Cultures were incubated overnight at 37°C (14–16 h), then terminated by centrifugation and removal of 200 \(\mu\)l supernatant into a separate tube. Both tubes were then counted in an LKB gamma counter for 5 minutes each. Controls included target cells alone in TCM for calculation of spontaneous chromium release, identical cultures freeze-thawed to determine maximum release, and mitogen-stimulated cultures. The mitogens used were PHA or con-A in optimal concentration for lymphocyte transformation.

**CALCULATION OF RESULTS**

Percentage chromium release was calculated as

\[ \% \text{ Cr release} = 100 \times \frac{2S}{S + R} \]

where \(S=\text{counts in supernatant} \)

\(R=\text{counts in residual volume.} \)

Specific cytotoxicity was calculated as

\[ \text{sp. cytotoxicity} = 100 \times \frac{T - S}{M - S} \]

where \(T=\text{mean }\%\text{ test Cr release of triplicates as calculated above,} \)

\(S=\text{mean }\%\text{ Cr release of 6 control cultures of target cells alone in TCM after overnight culture (spontaneous Cr release),} \)

\(M=\text{mean }\%\text{ Cr release of 6 control cultures of target cells alone in TCM after freezing and thawing 4 times and overnight culture (maximum release).} \)

**EVALUATION OF RESULTS**

Results were rejected if spontaneous chromium release exceeded 35% or if mitogen-induced cytotoxicity did not exceed spontaneous cytotoxicity. Mean standard error of the mean (SEM) for each group of values from individual assays was calculated and varied from 0.7 to 1.3, average 1.0. Results from a few triplicates with SEM greater than 2.5 were rejected. Unpaired and paired Student's \(t\) tests were used for statistical evaluation as indicated in the text.

**Results**

The results of cytotoxicity for PBM from rheumatoid and normal individuals against the 4 cell lines are shown in Fig. 1. There is no significant difference
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in spontaneous cytotoxicity between the rheumatoid patients and normal persons for any of the cell lines. It can be seen from Fig. 1 that there was a wide variation in cytotoxicity particularly for CCRF/CEM in both the rheumatoids and the normals. This scatter is due to variation between individuals in the figure, though there was also some variation in cytotoxicity by PBM from the same individual when tested on different occasions. Results for RPMI, RAJI, and Wil are more closely grouped than for CCRF with the exception of a few markedly higher results. Subjects who showed high cytotoxicity against one cell line tended to show high levels against the other cell lines, suggesting that this was not a specific phenomenon (results not shown).

The 2 highest values for spontaneous cytotoxicity against Wil were shown by patients with rheumatoid arthritis. One rheumatoid patient showing a value of 20.1 with Wil also had a high value of 59.8 against CCRF. The other extremely high value of 60.7 against Wil was determined solely against this cell line and cannot be compared with cytotoxicity to other cell lines simultaneously. However, it may well be anomalous, as this patient previously showed a value of 12.4 (not shown as excluded by high spontaneous chromium release). At the time of the high value the patient's clinical status had shown no obvious change and she was not suffering from an obvious viral infection. She had recently had a test injection of sodium aurothiomalate, but other patients on this therapy showed results within the normal range. Addition of sodium aurothiomalate up to 100 μg/ml to cultures in vitro had no significant effect on cytotoxicity (results for 7 RA subjects, 6 normal subjects not shown).

Discussion

No difference has been shown between the spontaneous cytotoxicity of peripheral blood mononuclear cells from patients with rheumatoid arthritis or from controls against the 4 cell lines tested. Though a positive result might have indicated a host response against lymphoblastoid antigens, the significance of a negative result cannot be so easily interpreted, while the mechanism by which SK cells recognise their targets is unknown. Thus, although rheumatoid patients do not preferentially kill cell lines positive for EBV genome, this cannot be taken as direct evidence that this agent could not play a role in the pathogenesis of RA, although other evidence against this hypothesis has recently been presented.17

There are a number of other factors to be considered. Although 2 groups were able to demonstrate increased killing of EBV-containing lymphoblastoid cell lines during active infectious mononucleosis using unfractinated peripheral blood leucocytes,18 10 a further group was able to demonstrate this effect only after removal of complements receptor bearing cells.20 Thus it is possible that a response of a sub-group of cells would be masked by nonspecific cytotoxicity in the experiments reported here. Interestingly, the increased cytotoxicity found...
during active EBV-induced infectious mononucleosis and thought to be a specific effect of circulating atypical T blast-like cells, has recently also been found in cytomegalovirus mononucleosis, again casting some doubt on the specificity of this finding. In both cases increased cytotoxicity was present only during actual infection and returned to normal levels during convalescence.

Of potential interest in this study were 2 rheumatoid patients showing high levels of cytotoxicity against Wil. However, one of these also showed a high level of cytotoxicity against CCRF, suggesting this might be a nonspecific effect, and the other value appears possibly anomalous in that the same patient had previously shown a value within the usual range, and no readily identifiable factors were associated with the second high value.

In general, results for rheumatoid patients in this study are similar to others previously reported for normal subjects. Thus the tumour cell line CCRF/CEM is relatively well killed, whereas Wil, derived from normal lymphocytes, is not significantly killed. RAJI is poorly killed but is known to be an exception to the rule that tumour cell lines are well killed in comparison to lines derived from normal lymphocytes. RPMI 8226 is an intermediate position. Thus, in studying cytotoxicity of unfractionated PBM from rheumatoids and normals against 4 lymphblastoid cell lines we have been unable to demonstrate any increase in cytotoxicity amongst the rheumatoids, either overall or against particular cell lines, and so, using this rather limited system, we have not been able to show preferential recognition of antigens on these cell lines by rheumatoids.

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