Specific (EMT6) and non-specific (WEHI-164) cytolytic activity by host cells infiltrating tumour spheroids

K.M. Wilson & E.M. Lord

Cancer Center and Department of Microbiology and Immunology, University of Rochester, Rochester, NY 14642, USA.

Summary The development of a serum-free, low-protein culture medium has allowed the detection of tumour-specific cytolytic cells in EMT6 immunized mice bearing EMT6 multicellular tumour spheroids. Spheroid associated (SAC) and peritoneal (PC) effector cells were specific to EMT6 as the target cell, not killing line 1, P815 or RIF-1. The natural killer (NK) cell sensitive target YAC-1 was also not lysed by SAC or PC, indicating undetectable levels of NK cells present within infiltrated spheroids. In contrast, high levels of cytolytic activity were present in SAC, PC and spleen cells against WEHI-164, a line sensitive to natural cytotoxic (NC) and macrophage mediated killing. The EMT6 specific activity was mediated by Thy1+, Lyt2+ cells. The anti-WEHI-164 effector cell population was Thy1+, Lyt2+. The WEHI-164 killer cells were present in SAC and PC from unimmunized mice while the EMT6 specific effector cells were not. After separation of SAC and PC by size using centrifugal elutriation, anti-EMT6 activity was present only in the lymphocyte fraction while anti-WEHI-164 activity was enriched in the macrophage fraction.

The multicellular tumour spheroid (MTS) (Sutherland et al., 1971) is a useful tumour model because it has many of the three dimensional characteristics of a solid tumour (Sutherland & Durand, 1984; MacDonald & Sordat, 1980). However, unlike solid tumours, MTS are easily dissociated into single cells by a mild trypsinization without loss of cell viability or function. We have used this model previously to demonstrate anti-EMT6 cytolytic activity by splenoid associated cells (SAC) from immunized spheroid bearing mice (Lord, 1980; Lord & Burkhardt, 1984). This anti-EMT6 activity was higher than that of peritoneal (PC) and spleen cells, showing the importance of studying in situ host cells (Lord & Burkhardt, 1984). However, since the immunizing EMT6 tumour cells and tumour spheroids were grown in foetal calf serum containing media, it was unclear whether the observed cytotoxicity represented specific anti-tumour immunity.

Cytolytic effector cells can be generated in vitro as the result of culture with foetal bovine serum (FBS), or in vivo following injection with FBS alone without specific antigen (Golstein et al., 1978). The activation of these cytolytic cells has been attributed to both the antigenic and mitogenic nature of xenogeneic serum. Since FBS components adsorbed or bound onto the surface of cells are not readily removed by washing (Golstein et al., 1978; Kubota, 1984), immunization with tumour cells grown in vitro in serum containing media could potentially elicit non-specific effector cells. We have demonstrated the generation of effector cell populations that lyse not only EMT6 but also a number of other tumour cell targets by inoculation of animals with EMT6 cells cultured in FBS supplemented media and/or culture of spleen cells from tumour bearing mice in FBS supplemented media (Reynolds et al., manuscript in preparation). Development of a serum-free, low-protein medium (MAT/P) (Taupier et al., 1985; 1986) in our laboratory for the culture of tumour cells and multicellular tumour spheroids has allowed us to demonstrate specific anti-tumour immunity.

In situ antigen specific Thy1+, Lyt2+ cytotoxic T lymphocytes (CTL) can be active in anti-tumour responses (DeLustro & Haskell, 1978). Natural killer (NK) cells and natural cytotoxic (NC) cells have been shown to have spontaneous cytolytic activity against a wide variety of tumours (Lattime et al., 1982). NK cells can be detected in a 4h chromium release assay and kill NK-sensitive in vitro grown tumour lines such as the YAC-1 lymphoma (Lattime et al., 1983). Except for a few animal studies (Ferry et al., 1984), NK activity within tumours has not been observed (Eremin et al., 1981; Totteman et al., 1980). NC activity is against some different targets such as the WEHI-164 sarcoma and is mediated by Thy1- cells (Lattime et al., 1982; 1983). Recently it has been shown that NC activity may be mediated by tumour necrosis factor (TNF) (Ortaldo et al., 1986). TNF has also been implicated in the lysis of WEHI-164 by mouse resident and peptone-induced peritoneal cells and human fresh, cultured and LPS stimulated monocytes (Colotta et al., 1986; Austgulen et al., 1986; Chen et al., 1985; Kildahl-Anderson et al., 1986; McKinnon, 1986; McKinnon et al., 1986). Since macrophages are present in high numbers in EMT6 solid tumours and MTS (Lord & Keng, 1984; Lord & Burkhardt, 1984), the non-specific cytotoxicity observed in the EMT6 system against WEHI-164 could be mediated by macrophages via TNF.

The objective of this research was to demonstrate specific anti-tumour cytolytic activity by in situ host cells in the absence of xenogeneic serum. We examined the specificity of anti-tumour cytolytic activity and characterized the effector cells responsible for anti-EMT6 and anti-WEHI-164 activity.

Materials and methods

Mice

BALB/cByJ female mice were purchased from the Jackson Laboratories, Bar Harbor, ME and used at 10–20 weeks of age.

Tissue Culture Medium

Complete MAT/P (CMP), a serum-free, low-protein medium (Taupier et al., 1985; 1986), was used for all experiments. Proteins used were insulin (0.005 mg/ml-1) and transferrin (0.005 mg/ml-1). All media were stored and used under yellow light. Tumour cells lines were adapted to serum-free medium by first reducing FBS concentration from 10% to 2% in MAT/P before using serum-free conditions.

Tumour system

EMT6 is a mammary sarcoma which arose spontaneously in a BALB/c mouse and was subsequently adapted for tissue culture (Rockwell et al., 1972). The subline designated EMT6/Ro (Rochester) was used for the experiments reported here. All cell lines were tested periodically for mycoplasmal contamination by Mycotrim (Hanna Media Inc., Berkeley, CA). EMT6/Ro was negative for retrovirus and 20 other
mouse viruses as tested by the MAP test (M.A. Bioproducts, Rockville, MD).

YAC-1, a lymphoma, was induced by Moloney virus in an A/Sn mouse (Cikes et al., 1973). WEHI-164, a fibrosarcoma, was induced by methylcholanthrene in a BALB/c mouse (Rollinghoff & Warner, 1973). RIF-1 (Twentyman et al., 1980), a fibrosarcoma, was induced by radiation in a C3H mouse. P815 (Dunn & Potter, 1957), a mastocytoma, was induced by methylcholanthrene in a DBA/2 mouse. Line-1 (Yuhas et al., 1974), a lung carcinoma, arose spontaneously in a BALB/c mouse.

Multicellular tumour spheroids

Multicellular tumour spheroids (MTS) were grown in vitro as previously described (Lord et al., 1979). Spheroids were initiated in vitro by placing 5 x 10³ EMT6 tumour cells into 100 mm non-tissue culture petri dishes (Labtek). After 4-5 days the small spheroids were transferred to spinner flasks containing 150 ml MAT/P and medium replenished daily. MTS of 600-800 μm diameter, obtained after ~2 weeks of growth, were used for all experiments.

Spheroid implantation and recovery

MTS (60/mouse) were implanted into the peritoneal cavity using an 18-gauge needle as previously reported (Lord et al., 1979). After 4-6 days spheroids were recovered by flushing the peritoneal cavity with a balanced salt solution (BSS) containing sodium heparin (Gibco) at 5 U ml⁻¹. The spheroids were separated from the peritoneal cells by unit gravity sedimentation. Spleens from the same animals were gently teased into single cell suspensions and washed with BSS. Spheroids were washed three times with BSS and dissociated into single cell suspensions with 0.05% trypsin for two 10 min periods at 37°C. Spheroid associated cells (SAC) were washed three times with cold BSS to remove trypsin.

Immunization

Mice were immunized 10-14 days prior to spheroid implantation with an i.p. injection of 5 x 10⁹ washed, irradiated (50 Gy) EMT6 cells from in vivo monolayer culture.

Cytotoxicity assay

An 18h chromium release assay was used to measure cytolytic ability of SAC, PC and spleen cells as previously described (Lord & Burkhardt, 1984a; Wilson & Lord, 1986). Two-fold dilutions of effector cells were prepared in 96 well flat bottomed plates. Ten thousand ⁵¹⁶Cr labelled exponentially growing EMT6 target cells were added to each well. The plates were incubated 18h at 37°C, then 0.1 ml supernatant removed and counted for radioactivity. Spontaneous release was measured by incubating targets with media alone. Complete release was determined by adding a 1:20 dilution of ZAP-isoton (Coulter Electronics) to targets. Four hour assays were identical, with the exception that V bottom plates were used and spun at 200 r.p.m. for 2 min at room temperature after addition of cells. Percent specific lysis (%SSL) was calculated from release of radiolabel from target cells, according to the following formula:

\[
\text{SL} = \frac{\text{(experimental mean–spontaneous mean)}}{\text{(complete mean–spontaneous mean)}}
\]

Morphological analysis

Slides were prepared using a Shandon–Elliot cytocentrifuge. The slides were air-dried, stained with the Gugol-blue (Wright-Giemsa) method and differential counts made on 300-500 cells per slide.

Antibody plus complement treatment

Treatment with cytotoxic monoclonal antibodies (ascites) plus complement was used to remove cell subpopulations prior to chromium release assay. Anti-Thy.2 was produced by H013.4 (Marshak-Rothstein et al., 1979), and anti-Lyt2.2 by H02.2 (Raulet et al., 1980). Cells at 1 x 10⁶ ml⁻¹ were suspended in the appropriate dilution (determined by titration) of antibody and placed on ice for 30 min. The cells were then centrifuged, resuspended in the appropriate dilution (determined by titration) of rabbit complement (CedarLane), and incubated at 37°C for 30 min. The cells were then washed, resuspended in medium and counted.

Centrifugal elutriation

A centrifugal elutriation procedure was utilized to separate subpopulations of SAC and PC (Lord & Keng, 1984). Single cell suspensions from implanted spheroids were elutriated in media containing 10% newborn calf serum at 4°C. The flow rate was raised from 25 to 41 ml min⁻¹ while the rotor speed was decreased in increments from 4000 to 0 r.p.m. and fractions collected at each interval. Lymphocyte fractions had a mean cell volume of 300 μm while macrophage fractions contained cells ranging from 700-1500 μm.

Results

Cytolytic activity in the absence of foetal bovine serum

We have previously determined that spheroid associated cells (SAC) recovered from mice immunized with irradiated EMT6 cells and injected with EMT6 multicellular tumour spheroids had cytolytic activity against radiolabelled EMT6 cells (Lord and Burkhardt, 1984). It was a concern that this activity might be the result of bovine serum components adhering to the immunizing cells since xenogeneic serum has been show to elicit cytotoxic cells (Golstein et al., 1978; Reynolds et al., manuscript in preparation). To overcome this potential problem, a serum-free, low-protein defined media (MAT/P) was used for growth of immunizing cells and spheroids, and in the chromium release assay. The added proteins, insulin and transferrin, do not interfere with generation of specific immunity (Reynolds et al., manuscript in preparation). Growth kinetics (size, plating efficiency and number clonogenic cells per spheroid) of tumour spheroids implanted in immunized and unprimed mice using serum-free conditions were similar to those previously published using FBS containing media (Lord & Burkhardt 1984; data not shown).

An 18h chromium release assay was performed using spheroid associated cells (SAC), peritoneal cells (PC) and spleen cells from immunized spheroid bearing mice as effectors, and monolayer EMT6 as radiolabelled targets. SAC were found to possess the greatest cytolytic ability against EMT6 (Figure 1). The PC also demonstrated cytolytic activity although to a slightly lesser extent. Spleen cells from these mice had little or no cytolytic ability. Thus the most cells infiltrating tumour spheroids were cytolytic in the absence of FBS, and had greater activity than those cells of the periphery.

Specificity of in situ and peritoneal effector cells

Lysis of EMT6 target cells is observed only in animals immunized with EMT6: no anti-EMT6 activity was detected from SAC when mice were immunized with a different tumour (Lord & Burkhardt, 1984). However, we wished to determine whether the cytolytic effector cells were specific for EMT6 target cells only. To determine the target specificity of SAC and PC, a panel of radiolabelled targets was used in a chromium release assay. RIF-1 (a C3H radiation induced fibrosarcoma) and line 1 (a BALB/c lung carcinoma) were not lysed by SAC and PC effector cells (Table
Table I Target lysis specificity of SAC, PC: % specific lysis at various effector to target ratios

| Target       | SAC       | PC       |
|--------------|-----------|----------|
|              | 35:1      | 18:1     | 9:1      | 100:1 | 50:1 | 25:1 |
| Experiment 1 | 18 hour chromium release assay | | | | |
| EMT6         | 51.1      | 52.0     | 38.8     | 51.5  | 34.6 | 24.5 |
| Line-1       | 0         | 0        | 6.2      | 3.2   | 0    | 0    |
| P815         | 39.1      | 7.1      | 11.9     | 28.1  | 2.8  | 0    |
| Experiment 2 | 18 hour chromium release assay | | | | |
| EMT6         | 24.4      | 21.6     | 10.9     | 23.0  | 18.9 | 9.1  |
| RIF-1        | 1.5       | 5.8      | 6.6      | 9.8   | 3.0  | 3.6  |
| Experiment 3 | 18 hour chromium release assay | | | | |
| EMT6         | 66.8      | 54.9     | 43.5     | 61.4  | 53.5 | 36.0 |
| WEHI-164     | 17.3      | 20.0     | 31.8     | 35.1  | 44.4 | 36.6 |
| Experiment 3 | 4 hour chromium release assay | | | | |
| EMT6         | 18.8      | 11.7     | 6.4      | 16.3  | 9.3  | 3.9  |
| WEHI-164     | 4.2       | 4.3      | 3.1      | 7.6   | 4.3  | 2.5  |
| YAC-1        | 3.9       | 3.0      | 2.5      | 19.4  | 16.3 | 10.5 |

I). P815 (a DBA/2 mastocytoma) was not killed except at the highest effector to target ratio which may have been influenced by crowded culture conditions. YAC-1, an A/Sn lymphoma and NK sensitive line, was not killed by SAC in a 4 h chromium release assay although some lysis by PC was present. A 4 h assay was used for this target since non-specific lysis by radiolabelled YAC-1 cells was too great at 18 h to yield meaningful data. WEHI-164, an NC and macrophage but not NK sensitive line (Lattime et al., 1982; 1983; Chen et al., 1985; Ortaldo et al., 1986) was killed by both the SAC and PC effector cells.

Characterization of the specific (anti-EMT6) and non-specific (anti-WEHI-164) cytolytic activity

The anti-WEHI-164 killing was very high, especially by SAC and PC from immunized mice (Figure 2). This killing was consistently higher than the anti-EMT6 activity at the lower effector to target ratios (Figures 1 and 2).

The anti-WEHI-164 but not the anti-EMT6 activity was present in spleens from splenoid bearing mice (Figures 1 and 2). The anti-WEHI effector cell was also demonstrable in SAC and PC from unprimed mice while the anti-EMT6 cell was not (Figure 3).

It was of interest to determine whether the anti-EMT6 and anti-WEHI-164 activities were carried out by the same or different cell populations. To determine the phenotype of the anti-EMT6 and anti-WEHI-164 effector cells, SAC and PC were treated with either anti-Thyl or Lyt2 plus complement prior to the chromium release assay. Antibody plus complement treatment removed the anti-EMT6 activity. Thus Thyl+, Lyt2+ cells were responsible for the anti-EMT6 activity (Figure 4). Antibody plus complement treatment did not remove anti-WEHI-164 activity; therefore, the anti-WEHI-164 effector cells were Thyl+ , Lyt2+ (Figure 4). These data indicate that the EMT6 and WEHI-164 killers are different cell populations.

To further characterize the cells responsible for killing EMT6 and WEHI-164, centrifugal elutriation was used to

Figure 1 Comparison of in situ and peripheral cytolytic activity in the absence of foetal bovine serum. 18 h chromium release assay of SAC (△), PC (□) and spleen cells (●) from immunized MTS bearing mice against EMT6 target cells. The difference between SAC and PC cytolytic activity was significant at the 95% confidence interval by lytic units analysis.

Figure 2 TitratioHex essay of anti-WEHI-164 cytolytic activity. 18 h chromium release assay of SAC (△), PC(□) and spleen cells (●) against WEHI-164 targets.

Figure 3 Comparison of cytolytic activity against EMT6 and WEHI-164 in immunized spheroid bearing mice. 18 h chromium release assay of SAC (●) and PC (□) vs. EMT6 (A) and WEHI-164 (B).
A

B

C

D

Figure 4 Characterization of effector cell responsible for cytolytic activity. 18 h chromium release assay of SAC (A, B) and PC (C, D) against EMT6 (A, C) and WEHI-164 (B, D) targets. Cells were treated with media alone (○), complement alone (□), anti-Thyl+complement (○) or anti-Lyt2+complement (△) immediately prior to assay.

Table II Differential counts from elutriated fractions

| Fraction   | % Lymphocytes | % Macrophages | % Granulocytes | % Tumour cells |
|------------|---------------|---------------|----------------|----------------|
| SAC unsep. | 9.2 ± 0.5a    | 34.0 ± 7.2    | 8.8 ± 0.5      | 48.0 ± 6.7     |
| SACL       | 56.6 ± 7.2    | 14.4 ± 1.6    | 29.0 ± 8.8     | 0              |
| SACM       | 2.5 ± 1.5     | 70.4 ± 2.9    | 0.8 ± 0.2      | 26.3 ± 4.2     |
| PC unsep.  | 57.1 ± 4.9    | 34.7 ± 3.8    | 8.2 ± 1.0      | 0              |
| PCL        | 82.5 ± 10.0   | 6.4 ± 1.0     | 11.0 ± 11.0    | 0              |
| PCM        | 15.7 ± 6.3    | 79.6 ± 6.0    | 4.7 ± 0.4      | 0              |

*SEM.

Discussion

The multicellular tumour spheroid (MTS) is useful as a tumour model due to its tumour-like 3-dimensional structure and ease in manipulation (Sutherland et al., 1971; MacDonald & Sordat, 1980; Sutherland & Durand, 1984). This model is especially useful for examining the host cells which infiltrate tumours. These in situ host immune cells (macrophages, granulocytes, lymphocytes) are often present in different proportions and activity levels than those cells of the periphery (Lord, 1980; Lord & Burkhardt, 1984). The MTS model was used to characterize the infiltrating host cell population responsible for cytolytic activity against tumour cells and the target lysis specificity.

It was a concern that the cytolytic activity of SAC demonstrated in previous studies might be due to foetal bovine serum used to culture cells used in immunization and growing MTS. To rule out this possibility, a serum-free, low-protein medium was developed in our laboratory (Taupier et al., 1985; 1986) and used exclusively in these experiments. Cytolytic activity was present in infiltrated EMT6 spheroids from immunized syngeneic BALB/c mice (Figure 1). This activity was greater than that of peritoneal cells (PC) while spleen cells had very little activity.

EMT6 spheroids implanted into immunized syngeneic mice became infiltrated by macrophages, granulocytes and lymphocytes (Lord & Burkhardt, 1984; Table II). Removal of either Thyl" or Lyt2" cells abrogated activity (Figure 4). Thus the cell responsible for anti-EMT6 cytotoxicity was a Thyl", Lyt2" cytotoxic T lymphocyte.

The target cell specificity of SAC and PC was determined by utilizing a panel of radiolabelled target cells in the chromium release assay. The SAC effector cells failed to kill several tumour lines including YAC-1, an NK-sensitive line (Table I). This is in agreement with some others who have been unable to demonstrate NK cells within tumours (Eremin et al., 1981; Totterman et al., 1980). It was separate SAC and PC subpopulations based on size. Differential counts of the lymphocyte and macrophage fractions (Table II) indicated a 5 to 20 fold enrichment for one cell type over the other. Anti-EMT6 activity was present exclusively in the lymphocyte fraction (Figure 5). Anti-WEHI-164 activity was present in each fraction but greatest in the macrophage fraction (Figure 5).
surprising, however, that the SAC effector cells killed WEHI-164 to an even greater extent than EMT6 (Figures 1 and 2). These anti-WEHI-164 killers (Thyl-1, Lyt2-2) were a different population of cells than the anti-EMT6 effectors (Thyl-1, Lyt2-2) (Figure 4). This non-specific activity against WEHI-164 was also present in host infiltrated tumour spheroids of line 1, a progressive BALB/c lung carcinoma (data not shown). The anti-WEHI-164 killers were present in spleen cells, SAC and PC from unimmunized mice where anti-EMT6 activity was not evident (Figure 3).

The anti-WEHI-164 activity may have been mediated by NC cells (Lattime et al., 1982; 1983). Recently it has been shown that NC activity may be mediated by TNF, a macrophage derived factor (Ortaldo et al., 1986). Since macrophages are present in high numbers in situ in the EMT6 system (Lord & Burkhartd, 1984; Lord & Keng, 1984; Table II) it is possible that this activity was mediated by a macrophage secreted cytokinin such as TNF (Kildahl-Anderson, et al., 1986; Chen et al., 1985; Colotta et al., 1986; Austgulen et al., 1986). In support of this hypothesis we found that anti-WEHI-164 activity was greater in fractions of effector cells enriched for macrophages by centrifugal elutriation than in fractions enriched for lymphocytes (Figure 5). These results indicate that while a lymphocyte population is responsible for the specific anti-EMT6 activity, the anti-WEHI-164 activity may be mediated by macrophages. This activity is present at very low effector to target ratios and is very high in the in situ host cell population. Preliminary data suggest that this activity is carried out by a secreted factor (data not shown).

It is interesting that while anti-EMT6 activity was exclusively in the lymphocyte fraction, anti-WEHI-164 activity was present to some extent in all fractions although it was highest in the macrophage fraction. This may be due to factor production by the small number of contaminating macrophages over the 18 h assay period, since the activity seen is high even at very low effector to target ratios. It is also interesting that EMT6 grows progressively as a solid tumour or tumour spheroids in unimmunized mice (data not shown). This progressive tumour growth occurs despite the presence of macrophages capable of cytolytic activity. These results indicate that EMT6 is insensitive to this macrophage mediated killing, but does not rule out possible cytoplastic effects.

We have demonstrated direct cytotoxicity of EMT6 by host cells recovered from MTS in a serum-free system. These anti-EMT6 effectors cells were Thyl-1, Lyt2-2 and specific for EMT6 targets. Thyl-1, Lyt2-2 cells, most likely macrophages, which mediated WEHI-164 killing were also present in SAC and PC from unimmunized as well as immunized spheroid bearing mice. Additional experiments are in progress to further characterize the WEHI-164 killing and its significance in anti-tumour immunity.

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Figure 5 Separation of effector cells by centrifugal elutriation. 18 h chromium release assay of SAC (A, B) and PC (C, D) against EMT6 (A, C) and WEHI-164 (B, D) targets. Cells were separated by centrifugal elutriation into lymphocyte (○) and macrophage (●) enriched fractions prior to assay.
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