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IN VITRO GENERATION OF HUMAN CYTOTOXIC LYMPHOCYTES BY VIRUS
Viral Glycoproteins Induce Nonspecific Cell-mediated Cytotoxicity without Release of Interferon*

By PAOLO CASALI, J. G. PATRICK SISSONS, MICHAEL J. BUCHMEIER, AND MICHAEL B. A. OLDSTONE

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The outcome of a virus infection represents an imbalance toward either the infecting agent's virulence or the host's ability to limit virus spread. Limiting infection may be a function of the immune system and take the form of a specific response and/or result from a nonimmune (nonspecific) response. Recently, considerable attention has focused on nonspecific host responses because of their potential to halt or reduce viral replication early in infection before the generation of either specific antibodies or cytotoxic T lymphocytes. Examples of early antiviral mechanisms are serum complement that can lyse some virions (1, 2) and bind to infected cells (3-5) in the absence of antibodies, natural killer (NK) cells (6) that lyse both infected and uninfected target cells (7, 8), and interferon, which can inhibit viral replication (9-12) as well as induce new or enhance ongoing NK cell activity (13-16).

The induction or enhancement of nonspecific cell-mediated cytotoxicity (CMC) (also called spontaneous CMC, natural CMC, or NK activity) by viruses and/or interferon has recently received much attention (6-8, 17-22). However, little is known about the role of viral polypeptides in induction or enhancement of this CMC. Early in infection, viral glycoprotein may be expressed on the surface of infected cells (23) before its release either into the fluid phase in soluble form or its incorporation into virions. For this reason, we studied the role of and mechanism(s) whereby soluble viral glycoproteins, viral glycoproteins inserted into artificial membranes (virosomes), and whole virions mediate nonspecific CMC.

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This research was supported by grants NS-12428, AI-07007, and AI-16102 from the U.S. Public Health Service and an investigator award by the medical group of Scripps Clinic and Research Foundation to J. G. P. S. and P. C. This work was done during the tenure of an Established Investigatorship from the American Heart Association to M. J. B.

1 Abbreviations used in this paper: ADCC, antibody-dependent cell-mediated cytotoxicity; CMC, cell-mediated cytotoxicity; CTL, cytotoxic T lymphocytes; E, erythrocytes; EA, antibody-coated erythrocytes; EYPC, egg yolk phosphatidylcholine; FBS, fetal bovine serum; F, measles virus fusion glycoprotein; HA, measles virus hemagglutinin; K cells, killer cells; LCMV, lymphocytic choriomeningitis virus; LU, lytic units; MEM, minimum essential medium; NK, natural killer; NP-40, Nonidet P-40; octylglucoside, octyl-β-D-glycopyranoside; ORBC, ox erythrocytes; PBL, peripheral blood lymphocytes; PBS, phosphate-buffered saline; RDMC, rhabdomyosarcoma cells; SDS-PAGE, sodium dodecyl sulfate-polyacrylamide gel electrophoresis; SRBC, sheep erythrocytes; SSPE, subacute sclerosing panencephalitis; VSV, vesicular stomatitis virus.
We found that purified measles virus glycoproteins in a soluble form or inserted into phospholipid vesicles induced significant cytotoxic activity in peripheral blood lymphocytes (PBL) within 2–4 h of incubation. This CMC was expressed as lysis of autologous or heterologous cells and was not associated with the release of interferon in culture fluids. In contrast, whole virions required >8 h to generate the same degree of cytotoxic activity. This activity was associated with interferon release in the culture fluids. Glycoproteins purified from lymphocytic choriomeningitis virus (LCMV) duplicated this result, suggesting that these findings may be a generalized phenomenon.

Materials and Methods

Virus Source, Purification of Viral Polypeptides, and Their Integration in Artificial Lipid Membranes. Measles virus and LCMV were used in this study. The source of measles virus, its cloning, passage history, method of propagation handling, and titration have been reported by this laboratory (3, 24, 25). To label virus, infected HeLa cells were incubated for 24 h in leucine-free Dulbecco's modified Eagle's minimum essential medium (MEM) supplemented with 10% heat-inactivated dialyzed fetal bovine serum (FBS) (Flow Laboratories, Rockville, Md.), 1% glutamine, penicillin (100 IU/ml), streptomycin (100 µg/ml), and 6.25 µCi/ml of [3H]leucine (Amersham/Searle Corp., Arlington Heights, Ill.). Culture fluid containing measles virus was freed from debris by an initial centrifugation at 1,500 g for 30 min at 4°C and the virus was then purified as described by this laboratory (24, 25). To prepare hemagglutinin (HA) and fusion (F) glycoproteins, purified virus solubilized in 0.15 M NaCl, 0.01 M Tris-HCl buffer, (Tris-NaCl buffer), pH 8.0, containing 2% Nonidet P-40 (NP-40) (BDH Chemicals, Poole, England) was applied to a column of Sepharose beads bearing covalently linked Lens culinaris agglutinin (LCH Gel; E. Y. Laboratories Inc., San Mateo, Calif.) (25). After adsorption of applied material, the column was washed with Tris-NaCl buffer containing 0.1% NP-40, which was later replaced with Tris-NaCl buffer containing 30 mM octyl-β-D-glucopyranoside (octyl-glucoside; Calbiochem, La Jolla, Calif.). Solid-phase-bound measles virus glycoproteins were then eluted using Tris-NaCl buffer containing 30 mM octylglucoside and 0.1 M α-methylmannoside. Protein concentration was determined by the Lowry method (26). As a control, a Sepharose-lentil-lectin column was run under similar conditions except that no solubilized virus was added. The fraction eluted from this column by α-methyl-mannoside was found to be protein free. The source, cloning, preparation, and handling of LCMV have been described (27). LCMV glycoproteins were purified by affinity chromatography with two serial passages through a lentil-lectin column using 2% sodium cholate as a detergent. The purity of viral glycoproteins preparations were assayed by electrophoresis on sodium dodecyl sulfate-polyacrylamide gels (SDS-PAGE) (25). Purified glycoproteins and the equivalent fraction eluted from the control column were freed of detergent by exhaustive dialysis at 4°C against phosphate-buffered saline (PBS). The integration of purified measles virus glycoproteins into lipid vesicles was carried out using egg yolk phosphatidylcholine (EYPC) (Sigma Chemical Co., St. Louis, Mo.) as reported (25). Traces of 3H-labeled measles virus glycoproteins and of 14C-labeled EYPC were used to check the integration of glycoproteins into the membranes by flotation gradient ultracentrifugation analysis (25).

Immunochemical Reagents and SDS-PAGE. Antibody to measles virus glycoproteins was obtained from the serum of a patient with subacute sclerosing panencephalitis (SSPE) and accounted for 10% of the total immunoglobulin (28). IgG antibodies and their F(ab')2 fragments were prepared as reported (3). Serum from a healthy donor (SH) with no detectable anti-measles virus antibodies, as assayed by neutralization, fluorescent binding to virus-infected cells, and immune precipitation tests with 35S-labeled measles virus (24, 29), was used as a source of IgG or F(ab')2 free of anti-measles viral antibodies. SDS-PAGE analysis of viral polypeptides was performed as described by Laemmli (30) and modified in this laboratory (24, 25).

Cytotoxicity Studies. PBL obtained from healthy measles immune and nonimmune subjects were used as effector cells in all cytotoxicity assays. Lymphocytes were obtained from heparin-
induced peripheral blood centrifugated on a Ficoll-Hygapaque gradient (31) and freed of adherent cells by incubation for 1 h at 37°C with 5% CO₂ in a tissue culture flask (175 cm²; Falcon Labware, Oxnard, Calif.). To obtain lymphocyte subsets, PBL were first incubated with neuraminidase- (Gibco Laboratories, Grand Island Biological Co., Grand Island, N. Y.) treated sheep erythrocytes (SRBC) (32) and segregated into erythrocyte-rosette forming (E⁺) cells and non-E-rosette forming (E⁻) cells by application to a Ficoll-Hygapaque gradient. Lymphocytes with (C3⁺) and without (C3⁻) receptors for C3 were separated by using a column of Sepharose 4B beads (Pharmacia Fine Chemicals, Piscataway, N. J.) bearing activated human C3 molecules, as previously described (33). Lymphocytes bearing receptors for the Fc portion of IgG (Fc-positive cells) were removed from PBL by using a monolayer made of ox erythrocytes (ORBC) coated with rabbit anti-ORBC IgG. Tissue culture flasks (T-25; Falcon, Labware) previously treated with PBS containing 100 μg/ml of poly-L-lysine (70,000 mol wt) (Sigma Chemical Co.) were coated with ORBC during a 2-h incubation at room temperature. After washings, bound ORBC were allowed to react for 45 min at 37°C with an excess of rabbit anti-ORBC IgG. The erythrocyte antibody (EA) monolayer was then used to absorb Fc-positive cells by incubation for 1 h at 37°C. Under similar conditions, PBL were applied to an E monolayer (prepared as described for the EA monolayer, but without antibodies) to determine their degree of nonspecific binding.

Cell lines used as targets in this study were: adult human skin fibroblasts (for autologous cytotoxicity assays) and human fetal skin fibroblasts established in culture in our laboratory, K562 cells, rhabdomyosarcoma cells (RDMC; from the Wistar Institute) and Molt-4 cells. Fibroblasts and RDMC were cultured in MEM supplemented with 1% glutamine, 10% FBS, and antibiotics; K562 and Molt-4 cells were cultured in RPMI 1640 medium supplemented with 10% FBS, 1% glutamine, and antibiotics.

Cytotoxicity assays were carried out in 96-well plates (Costar, Cambridge, Mass.) using ⁵¹Cr-labeled target cells as described (28) with slight modifications. Specific lysis was calculated by the following formula: 

\[
\text{percent specific release} = \frac{\text{Sam} - \text{Spo}}{\text{Max} - \text{Spo}} \times 100.
\]

In most experiments, the effector to target cell ratio used was 100:1. When three effector to target cells ratios were used (100:1, 50:1, and 25:1), specific cytotoxicity was computed in lytic units (LU) over the number of effector cells by a modification of the Von Krogh equation (34). One LU is expressed as the number of lymphocytes required to lyse 50% of the target cells during the incubation time specified. All test cultures and controls were run in triplicate. Most often fibroblasts were subject to an 18-h assay; spontaneous release did not exceed 30%. Kinetic experiments with ⁵¹Cr-labeled fibroblasts as targets were designed as follows: at time 0, four plates containing identical reagents were set up and incubated at 37°C in 5% CO₂. Each plate included samples for analysis of both spontaneous and maximal release from ⁵¹Cr-labeled targets. At varying time-points, 200 μl of supernatant fluid was harvested from each well, 100 μl for determining ⁵¹Cr release and 100 μl for titrating interferon-like activity.

Interferon Assay. Interferon activity of supernatant fluids was titered as their ability to inhibit the cytopathic effects of vesicular stomatitis virus (VSV) on a monolayer of Vero cells as described by Dahl (35) with minor modifications. All titrations were carried out in triplicate and interferon units were expressed as the reciprocal of the dilution inhibiting cytopathic effects by 50%. Each assay included Vero cells treated with a reference interferon preparation (National Institutes of Health Reference Interferon G-023-901-5271). Under the conditions used, one interferon unit derived in experiments was equivalent to one reference unit of NIH Human Reference Interferon.

Results

Soluble Measles Virus Glycoproteins Induced CMC. We first determined whether soluble measles virus glycoproteins and measles virions could induce PBL taken from several
human donors to kill heterologous and/or autologous fibroblast targets. Either 600 ng of purified measles virus glycoproteins, 3,000 ng of untreated or ultraviolet light (UV)-inactivated measles virions, or PBS was mixed with $10^5$, $5 \times 10^5$, or $2.5 \times 10^5$ PBL and $^{51}$Cr-labeled target cells, and the specific $^{51}$Cr release was determined over an 18-h period. Table I shows that PBL from all nine individuals studied mediated cytotoxicity. PBL incubated with untreated or UV-inactivated virions released significantly more $^{51}$Cr from both autologous and heterologous targets than did PBL incubated in PBS ($P < 0.02$). Soluble measles virus glycoproteins induced as much or more killing by PBL as whole virions. One of the nine individuals, SH, had no history of measles virus infection and no antibody to measles, but her PBL were as effective as PBL from immune adults in inducing CMC after exposure to measles virus glycoproteins. Results were similar when PBL from these subjects were incubated with measles virus glycoproteins and tested against Molt-4, K-562, or RDMC target cells in a 4-h $^{51}$Cr release assay (data not shown). In contrast, incubation of these targets with virus glycoproteins in the absence of PBL did not yield any significant lysis. Additionally, the eluted fraction from a control Sepharose lentil-lectin column, devoid of measles virus glycoproteins, also failed to induce CMC.

We next determined the amounts of soluble measles virus glycoproteins required to cause PBL killing of target cells and the kinetics of lysis. Equivalent amounts of free glycoproteins and of glycoproteins associated with measles virions were studied for their induction of CMC. This was accomplished by using virions at protein concentration five times higher than that of soluble measles virus glycoproteins. Previous studies with lentil-lectin affinity chromatography indicated that glycoproteins accounted for ~20% of the whole virion protein (25). By establishing a dose-response curve, we noted a linear relationship between the amount of glycoproteins present and the percentage of specific $^{51}$Cr released from autologous fibroblasts in an 18-h assay (Fig. 1). As little as 100 ng of virus glycoproteins/$10^6$ PBL resulted in CMC.

### Table I

<table>
<thead>
<tr>
<th>Donor</th>
<th>PB only</th>
<th>Virions</th>
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<td>Autologous</td>
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<tr>
<td>SH</td>
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</table>

25.9 ± 26.2 30 ± 22.2 122.6 ± 48.2 95.0 ± 31.0 116.8 ± 48.8 91.9 ± 29.1 181.9 ± 57.9 150.2 ± 57.4

* An 18-h $^{51}$Cr release assay was used, and the data are presented as LU/$10^6$ PBL. The release of $^{51}$Cr from target cells alone over this timed period did not exceed 30%.

§ SH was a donor with no clinical history of measles virus infection and no detectable circulating anti-measles virus antibodies. All other donors had detectable levels of anti-measles virus antibodies.

§§ Mean value ± 1 SD.
Cell Binding Requirements of Measles Virus Glycoproteins for CMC. Next we determined whether the binding of virus glycoproteins to the surfaces of PBL and/or target fibroblasts was sufficient to induce CMC. After $1 \times 10^8$ PBL or $1 \times 10^4 \text{Cr}$-labeled autologous fibroblasts were preincubated with varying amounts of virus glycoproteins for 2 h at 37°C, the cells were either washed extensively to remove free glycoproteins or were transferred unwashed with their culture fluid (containing soluble glycoproteins) to plates containing untreated $\text{Cr}^5$ fibroblasts and PBL, respectively. We found that PBL exposed to virus glycoproteins, washed, and then added to fibroblast targets induced CMC, but approximately six times more glycoproteins ($2,900 \text{ ng/10}^6 \text{ PBL}$) were needed to reach 50% lysis than when soluble viral glycoproteins were kept in the incubation fluids ($500 \text{ ng/10}^6 \text{ PBL}$) (Fig. 2A). Similar results occurred when whole virions were used. CMC was induced when $\text{Cr}^5$-labeled fibroblasts were incubated with viral glycoproteins, washed, and added to untreated PBL. 10 times more glycoproteins ($>10 \mu\text{g/10}^4 \text{ fibroblasts}$) were needed to achieve 50% of lysis than when the soluble glycoproteins were kept in the culture fluid ($1.0 \mu\text{g/10}^4 \text{ fibroblasts}$) (Fig. 2B). Radiolabeling studies showed that at $1.0 \mu\text{g/10}^6 \text{ PBL}$ or $10^4 \text{ fibroblasts}$, ~2% of offered measles virus glycoproteins became cell associated under the experimental conditions used. Hence, these studies indicated first that viral glycoproteins bound to fibroblasts could induce cytotoxic activity in unprimed PBL, and second, that as little as $10-20 \text{ ng}$ viral glycoprotein bound to $10^6 \text{ PBL}$ induced significant CMC.

Inhibition of Measles Virus Glycoprotein-induced CMC by Anti-Measles Virus Antibody. F(ab')$_2$ fragments from an IgG fraction containing high titers of antibodies to measles virus and an IgG fraction devoid of anti-measles virus antibodies were compared for their ability to inhibit measles virus glycoprotein-induced CMC. PBS or PBS containing F(ab')$_2$ antibody in decreasing amounts from 4,500 to 4.4 µg were mixed a fixed amount of either measles virus glycoproteins ($1.0 \mu\text{g}$) or whole measles virus ($5.0 \mu\text{g}$) and incubated for 1 h at 37°C. Each mixture was then added to PBL ($1 \mu\text{g}$ of glycoproteins/$10^6 \text{ cells}$) and $\text{Cr}^5$-labeled autologous fibroblasts, and the specific $\text{Cr}$ release was determined over 18 h (Fig. 3A). F(ab')$_2$ antibody to measles virus (562
FIG. 2. Binding of measles virus glycoproteins to PBL or fibroblasts and enhancement of CMC.
Panel A: resultant cytotoxicity against 51Cr-labeled autologous fibroblasts. PBL incubated with increasing amounts of viral glycoproteins were transferred with fluid-phase glycoproteins (○) or in the absence of soluble glycoproteins (□) into culture wells containing 51Cr-labeled heterologous fibroblasts. Panel B: 51Cr-labeled heterologous fibroblasts were incubated with increasing amounts of measles virus glycoproteins and then left in culture with glycoproteins (○) or glycoprotein containing medium was removed (□). Thereafter, untreated PBL were added. In both experiments, the effector to target cell ratio was 100:1 and the assay time was 18 h.

Next, the ability of F(ab')2 antiviral antibody to block the cytotoxicity of PBL that have been already exposed to virus glycoproteins was studied. Various amounts of glycoproteins (45–5,800 ng) were incubated with 10^6 PBL for 2 h at 37°C. The PBL were washed extensively and added to either PBS, PBS containing F(ab')2 antibody (600 µg/10^6 PBL) to measles virus, or F(ab')2 (600 µg/10^6 PBL) preparation without antibody to measles virus. The mixtures were incubated with 51Cr-labeled autologous
fibroblasts and the amount of $^{51}$Cr released over 18 h was determined. As seen in Fig. 3B, antibody to measles virus added to PBL previously exposed to virus glycoproteins did not abrogate their ability to lyse target cells.

**PBL Subsets Responsible for Viral Glycoproteins Induced CMC.** Positive and negative selection experiments to determine the effector PBL subsets involved in CMC were performed with PBL from two donors for each of the two separate experiments. Total PBL, $E^+$ cells, $E^-$ cells, $E^-C3^-$ cells, $Fc^+$ cells, and $Fc^-$ cells were assessed for the capacity to mediate CMC against $^{51}$Cr-labeled fibroblasts in the presence of PBS, PBS containing 900 ng of measles virus glycoproteins, or whole measles virions containing 4,500 ng of protein. As seen in Fig. 4, the $Fc^+$ PBL subset, not $Fc^-$ cells, exerted CMC. Assays of other markers in PBL subsets placed CMC activity both in $E^+$ and $E^-$ cells, but $E^-$ and $C3^-$ cells were the highest in activity. Hence, lymphocytes with and without E receptors mediate cytotoxicity induced by measles virus glycoproteins. The non-E-rosetting PBL mediating CMC are C3 receptor negative. However, all PBL mediating cytotoxicity induced by measles virus glycoproteins or by measles virus have Fc receptors.

**Segregation of CMC Induced by Soluble Virus Glycoproteins from That Induced by Virions.** Unexpected but significant differences in the time-course of CMC activity occurred when PBL were incubated with either soluble measles virus glycoprotein or untreated or UV-inactivated virions. As demonstrated in Fig. 5A and C, as soon as 4 h after adding 900 ng of soluble measles virus glycoproteins to $10^8$ PBL, significant lysis of $^{51}$Cr-labeled autologous fibroblasts occurred. In marked contrast, equivalent lysis of fibroblasts by PBL incubated with untreated or UV-inactivated virions (4,500 ng of protein content) required 8 h or more of incubation. This difference in the early induction of CMC by soluble virus glycoprotein as compared with the later induction by whole virions was repeatedly observed in all experiments using PBL from eight individuals. Maximal $^{51}$Cr release occurring at 18 h was similar whether the PBL were incubated with soluble glycoproteins or virions.

![Fig. 4. Subsets of PBL responsible for CMC induced by viral glycoproteins. PBL or various subsets were assayed in the presence of PBS (●), measles virus glycoproteins (□), or whole measles virus (■). Panel A: positive selection experiment. PBL, nonadherent cells; $E^+$, lymphocytes forming rosettes with SRBC; $E^-$, lymphocytes not forming rosettes with SRBC; $E^-C3^-$, lymphocytes not forming rosettes with SRBC and without surface receptor for C3. Panel B: negative selection experiment. $Fc^+$ and $Fc^-$ PBL with or without surface receptor for the Fc fragment of IgG, respectively. In all experiments, the target cells were heterologous fibroblasts and the assay time was 18 h. The data represented were obtained by PBL from two donors. PBL from two other donors performed similarly.](image-url)
To further segregate these differences, we questioned whether soluble viral glycoproteins or virions from measles virus induced interferon-like activity in the supernatant fluids by PBL. In Fig. 5, panels B and D show that interferon was released into the culture medium only when untreated or UV-inactivated virions were present in culture with PBL. Interferon production peaked between 8 and 12 h, after which no further increase was demonstrable up to 18 h. The material was interferon-like in that it inhibited VSV replication, lost activity after trypsin treatment, and was not pelleted by ultracentrifugation (100,000 g/h). In none of our experiments did soluble measles virus glycoproteins incubated with PBL induce release of interferon-like activity in the culture medium. These findings were consistently reproduced and noted in PBL from several individuals.

We next determined whether the supernatant fluids harvested from the PBL-soluble glycoprotein cultures or from the PBL-virions cultures were able to induce CMC in newly added PBL. For this experiment, $1 \times 10^6$ PBL were incubated for 1 h at 37°C with either soluble measles virus glycoproteins (1,200 ng), virions (6,000 ng protein content), or PBS, and were washed and added to $51$Cr-labeled or unlabeled autologous fibroblasts, or incubated without fibroblasts. After 6, 12, and 18 h of incubation, 150 μl of supernates from various cultures was UV-irradiated for 5 min and then added with untreated PBL and $51$Cr-labeled fibroblasts in a 6-h culture. The degree of lysis achieved in the primary cultures is represented in Fig. 6 A. Results shown in Fig. 6 B indicate that media from PBL treated with whole virus and incubated with or without fibroblasts consistently induced CMC. In contrast, media from PBL preincubated with purified glycoproteins or PBS only, either in the presence or absence of fibroblasts, failed to induce cytotoxic activity in newly added PBL.

**Induction of CMC by Measles Virus Glycoproteins Integrated into Lipid Bilayers.** We next determined whether these glycoproteins must be extracted from the viral envelope.
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and presented in a free form to induce CMC, as in the foregoings experiments, or
whether this activity is retained when the glycoproteins are integrated into a lipid
bilayer. First, different amounts of purified measles virus glycoproteins from 28 ng to
3,600 ng, free or inserted into lipid bilayers, were added to $10^6$ PBL and cultured with
$1 \times 10^4$ $^{51}$Cr-labeled autologous fibroblasts. The amount of $^{51}$Cr released over 18 h
was then determined. In a second study, $10^6$ PBL were incubated with 900 ng of
purified measles virus glycoproteins, either in soluble form or integrated into vesicles,
and $10^4$ $^{51}$Cr-labeled fibroblasts. The $^{51}$Cr release was measured after 4, 8, 12, and 18 h.

As shown in Fig. 7, measles virus glycoproteins inserted into artificial lipid mem-
branes enhanced CMC to the same extent as an equivalent amount of their soluble
counterparts (panel A), and with the same kinetics (panel B). Vesicles with integrated
glycoprotein spikes (virosomes) did not induce release of mediators with interferon-
like activity in the culture medium (data not shown). EYPC vesicles without the virus
glycoproteins, “naked vesicles,” did not induce any CMC against fibroblasts.

**Effect on CMC of Combining Measles Virus Glycoproteins and Virions.** To determine
whether the virus glycoproteins and virions had an additive effect in inducing CMC,
$10^6$ PBL were incubated with $10^4$ $^{51}$Cr-labeled fibroblasts and 1.2 $\mu$g of soluble measles
virus glycoproteins, virions of measles virus containing 6 $\mu$g of protein, or both
glycoproteins (1.2 $\mu$g) and virions (5 $\mu$g of protein). As seen in Fig. 8A, by 12 h after
incubation, the combined reagents (glycoprotein and virion) caused a lower CMC
value than that resulting from incubation with measles virus glycoproteins alone.
Results with UV-inactivated virions were similar to untreated virions. At 18 h, the
combined effect of the two reagents was similar to that of glycoproteins or virus alone.
The soluble measles virus glycoproteins did not interfere with the release of interferon

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**Fig. 6.** Recruitment of cytotoxic lymphocytes by fluids from cultures of PBL and measles virions
or PBL and soluble glycoproteins. Panel A: CMC by PBL preincubated with measles virus
glycoproteins (O), virions (C), or PBS (A), washed and cultured with $^{51}$Cr-labeled heterologous
fibroblasts for the indicated time. The effector to target cell ratio was 100:1. Panel B: CMC
induction by culture fluids from PBL and measles virions with (C) or without (O) fibroblasts;
culture fluids from PBL and glycoproteins with (O) or without (O) fibroblasts; culture fluids from
PBL incubated with PBS in the presence of fibroblasts (A). Culture fluids were harvested at
indicated time. The effector to target cell ratio was 100:1 and the assay time was 6 h.
activity induced by the virions (Fig. 8B), because this activity was equal in culture fluids containing virions alone or with virions mixed with the glycoproteins.

Soluble LCMV Glycoproteins Induced CMC. PBL from several of the individuals tested above also killed human fibroblasts after incubation with purified LCMV glycoproteins and were evaluated by the same criteria used for measles virus. Purified LCMV glycoproteins enhanced CMC in a dose-response fashion as did measles virus glycoproteins (Fig. 1) and LCMV glycoprotein bound to the PBL as shown for measles virus glycoproteins (Fig. 2). In fact, LCMV soluble glycoproteins and whole virions behaved like the measles virus components in all parameters cited in this paper. Specifically, LCMV glycoproteins induced CMC faster than did whole LCM virions, and combining LCMV soluble glycoproteins and whole LCM virions failed to produce a cumulative CMC effect. Additionally, only the virions, not purified glycoproteins, induced release of interferon by PBL. Further, purified LCMV nucleocapsids failed to induce CMC under similar experimental conditions.
Here we report a novel mechanism by which CMC can be initiated and expressed. Purified soluble measles virus HA and F glycoproteins alone or inserted into lipid bilayers induce or enhance human PBL killing of a variety of target cells including autologous and heterologous human fibroblasts, Molt-4, K-562, and RDMC cells. Studies of the kinetics of CMC induction, the absence of interferon in the culture fluids, and the inability of supernates to recruit cytotoxic activity from newly added PBL distinguish this type of nonspecific human CMC from the interferon-associated enhancement of NK cell activity (13, 17, 20). The segregation of various nonspecific CMC activities is outlined in Table II. Cells involved in these different effector mechanisms cannot yet be separated into discrete subsets according to conventional surface markers (Fig. 4).

The killing of target cells by PBL induced with soluble measles virus glycoproteins was efficient and occurred early. After first exposing PBL to glycoproteins and then transferring them to cultures of $^{51}$Cr-labeled targets, we found that as little as 10 ng of glycoprotein bound to PBL induced CMC. Further, kinetic studies showed that in most experiments, significant lysis (20–40% of total specific $^{51}$Cr release) occurred within 2–4 h of exposing target cells to PBL in the presence of viral glycoproteins. Virus glycoproteins inserted into lipid bilayers performed similarly, indicating that the activation of PBL was not dependent on the hydrophobic region of the glycoproteins. Previous electron microscopic and binding activity studies in this laboratory (25) have indicated that the hydrophobic parts of the measles virus glycoproteins are inserted into the vesicle's lipid bilayer and that the "spike projections" are arranged in patterns identical to those in virions. In contrast to this activity of free glycoproteins or glycoproteins inserted into lipid bilayers, measles virion-induced CMC in PBL takes three to four times longer to be expressed and kill comparable numbers and types of target cells. The dose-response curves for viral glycoproteins in soluble form, inserted into vesicles, or in the whole virion were equivalent in terms of CMC induction. However, the activity of soluble glycoproteins and virions combined was

<table>
<thead>
<tr>
<th>Effector*</th>
<th>Inducer</th>
<th>Lysis of targets at‡</th>
<th>Culture fluids</th>
<th>Anti-viral activity§</th>
<th>Capacity to induce new CMC</th>
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<tr>
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</table>
| PBL | Virions | Nil | Nil | + | ++++ | ++++ | ++++

* Fe⁺ lymphocytes in both E⁻ and E⁺ populations.
‡ This evaluation is based on data from multiple quantitative experiments carried out under conditions similar to those described in Fig. 6, using $^{51}$Cr-labeled autologous or heterologous fibroblasts as targets. Relative cytotoxic index with a value of 1.0 being maximum: ±, <0.1; +, 0.1–0.25; ++, 0.26–0.5; ++++, 0.51–0.75; ++++, 0.76–1.0.
§ Interferon activity as judged by inhibition of cytopathic effect of VSV on Vero cells.
not additive but was intermediate between that of the two tested individually. This suggests that the whole virion per se might hamper early CMC induction due to glycoproteins alone. It is likely that both reagents compete for the same receptors on the cell's surface. The fact that UV-inactivated virions perform in a manner similar to their untreated counterparts suggests that replication of virus is not needed to inhibit early glycoprotein-induced CMC.

Supernatant fluids from PBL cultured in the presence of soluble viral glycoproteins or glycoproteins inserted in a lipid bilayer with or without fibroblasts did not contain interferon or other factors able to induce unprimed lymphocytes to be cytotoxic. In contrast, culture fluids from PBL incubated with virions, containing an equivalent amount of viral glycoproteins on their surface, displayed antiviral activity and were able to recruit lymphocytes to be cytotoxic.

Other experiments also indicate that viral glycoproteins absorbed on target cells are sufficient to induce CMC. Furthermore, the initiation of CMC activity by PBL with measles virus glycoproteins and virions was specific in that F(ab')2 antibody to measles virus glycoprotein blocked the induction of CMC by first binding to the virus glycoproteins or virion. This prevented their binding to PBL. However, once the PBL had been exposed to glycoproteins, the antibody had no effect on inhibiting CMC activity, suggesting that glycoproteins, in part, directly activate lymphocytes to be cytotoxic. There are two recent studies concerning glycoproteins and activation of CMC in vitro. With PBL from humans, Härfast et al. (36) showed that mumps virus glycoproteins prepared by trypsinization of the virion induced CMC as did whole virions. With mouse PBL, an interferon-independent mechanism of activation of NK cells by alloantibodies with specificity for NK cells or by Helix pomatia lectin has been reported (37). Culture fluids from lymphocytes activated by these stimuli also failed to recruit new cytotoxic lymphocytes. Hence, the CMC induced in mouse lymphocytes by Helix pomatia lectin or alloantibodies to NK cells appears similar to that reported here in human lymphocytes by viral glycoproteins.

What could be the biologic implications of our findings? Cytotoxic lymphocytes play an important role in limiting virus infection and in surveillance against tumor cells, and by their lytic action also cause tissue injury that can harm the host. Experimental models showed that cytotoxic lymphocytes generated during a viral infection in vivo can be segregated into two main sets. One set is immunologically specific in terms of activation, recognition, and killing of target cells. Included within this set are specific cytotoxic T lymphocytes (CTL) and lymphocytes that kill through the recognition and binding of the Fc portion of specific anti-viral IgG (antibody-dependent cell-mediated cytotoxicity [ADCC]). In vivo studies indicate that specific CTL and IgG antibodies are not generated until 5–7 d after appropriate antigenic stimulation (38, 39). However, by this time the infecting virus can go through multiple cycles of replication (40).

The second set of CMC is generated earlier in vivo, usually within the first 24 h of infection (41), but lacks the fine target specificity shown by CTL and ADCC. NK cells are part of this second set of early, unspecific CMC. NK cells are generated in vitro within 6–12 h after exposure of PBL to virions, virus-infected, or tumor-bearing cells. NK cells segregate with the Fc receptor-bearing lymphocytes (42). In vivo and in vitro induction of NK activity is associated with the release of interferon by PBL. From the data in this paper, viral glycoprotein-induced CMC may constitute an
earlier line of defense, because its expression occurs already within 2-4 h after the exposure of lymphocytes to glycoproteins and interferon release is not required. Other experiments (P. Casali and M. B. A. Oldstone, manuscript in preparation) indicate that similar to purified free glycoproteins, fibroblasts expressing measles virus HA and F polypeptides on their surfaces induce non-interferon-associated CMC within 2-4 h of incubation with PBL.

Thus, these early cellular mechanisms of defense against virus infection, comprising interferon-dependent as well as interferon-independent CMC, could help to limit virus spread by removing those cells that would otherwise later release infectious virus. Uninfected bystander cells could be killed in the microenvironment where these cells are operating. The combined action of killer cells induced by virus glycoproteins, virions, and interferon may function to limit both the spread of virus and the numbers of infected cells in primary infections until specific CTL and antiviral antibodies are generated much later. Failure of these early nonspecific defense mechanisms might favor the virulence of the virus over the host’s immune system later in infection.

Our observations with two dissimilar RNA viruses, measles virus and LCMV, suggest that other viruses may perform similarly. Future work will determine both the generalities of these findings for other viruses and its biological significance in vivo.

Summary

Purified hemagglutinin and fusion glycoproteins of measles virus either in soluble form or inserted in artificial membranes bind to human peripheral blood lymphocytes and induce cell-mediated cytotoxicity (CMC) in a dose-response fashion. Both autologous and heterologous noninfected target cells are lysed in vitro. The expression of CMC is not inhibited by anti-measles virus antibody added to lymphocytes previously exposed to viral glycoproteins. The killer lymphocytes are Fc receptor positive, both erythrocyte-rosetting and non-erythrocyte-rosetting, as assessed by both positive and negative selection experiments.

The induction of nonspecific CMC by viral glycoproteins either in the soluble state or inserted into artificial membranes could be segregated from the CMC associated with whole virions. First, on kinetics studies, purified viral glycoproteins induced CMC more rapidly than did whole virions. Second, viral glycoprotein-produced response occurred in the absence of detectable release of interferon into the culture medium, whereas CMC activity due to whole virions was associated with interferon release. The fact that purified measles virus glycoproteins integrated into artificial membrane bilayers were as efficient as their soluble counterparts in inducing CMC suggests that the hydrophobic portion of the glycoproteins was not involved in the induction and expression of the lytic activity. Purified glycoproteins from lymphocytic choriomeningitis virus behave similarly, although this virus is unrelated to measles virus. It is inferred that interferon-independent CMC induced by viral glycoproteins might account for some of the biological reactions occurring early in the control of a viral infection.

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