CHARACTERISTICS OF NATURAL KILLER CELLS IN THE MURINE INTESTINAL EPITHELIUM AND LAMINA PROPRIA

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Morphologic and functional similarities between the lymphoid tissues of the gut and lung have shown that a common mucosal immune system is involved in the protection of mucosal sites (1). Several studies have demonstrated the relevance of antibody, particularly IgA, in mucosal immunity. However, less attention has been directed to the role of cell-mediated immunity. The recent development of methods for the isolation of mucosal lymphoid cells (2-7) has facilitated the analysis of cytotoxic functions of lymphocytes from intestine and lung. Thus, it has been shown that guinea pig and human lymphocytes from the intestinal mucosa exert mitogen-induced, antibody-dependent, and spontaneous cellular cytotoxicity (6-8). Spontaneous cytotoxic activity of cells from the mouse lung (9) and gut (10) has many characteristics of natural killer (NK)1 cytotoxicity.

The main effector cells of NK activity in human (11), rat (12), and mouse (13) are large granular lymphocytes (LGL) with a high cytoplasmic/nuclear ratio and azurophilic granules in the cytoplasm. Because large lymphocytes with cytoplasmic granules are abundant in the mammalian intestinal mucosa (2), we postulated that gut granulated lymphocytes (gGL) might be the NK effector cells at that site (10). Thus, we initiated a study to clarify the relationship between gGL and gut NK cytotoxicity. In an attempt to define the phenotype of gut NK cells and their lineage, experiments with sera against various surface markers were performed. Moreover, splenic and gut NK activity were compared in C57BL/6-bg/bg (beige) mice, a mouse strain with very low NK activity (14), and in WBB6F1-W/Wv mice, a hybrid mouse deficient in hemopoietic stem cells and mast cell precursors (15). The results obtained in this and previous studies (10) suggest that gut NK cells in the mouse are lymphocytes of the T cell lineage, similar but not identical to splenic NK cells.

Materials and Methods

Mice. Inbred CBA/J mice of both sexes were obtained from The Jackson Laboratory, Bar Harbor, ME. C57BL/6-bg/bg (beige) mice were bred in McMaster University from heter-

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§ Supported by grants from the Medical Research Council of Canada.

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Abbreviations used in this paper: A/T, attacker/target ratio; C, complement; gGL, gut granular lymphocytes; HBSS, Hanks' balanced salt solution; IEL, intraepithelial lymphocytes; LGL, large granular lymphocytes; LP, lamina propria; LPL, LP lymphocytes; LU, lytic units; NK, natural killer.

J. Exp. Med. © The Rockefeller University Press • 0022-1007/82/06/1785/12 $1.00 1785

Volume 155 June 1982 1785-1796
ozygotes purchased from The Jackson Laboratory. Similarly, WBB6F1-W/W mice were bred in McMaster University from WB/ReJ-W/+ and C57BL/6J-W/+ parental stocks purchased from The Jackson Laboratory. In this study, littermates of W/W mice included WBB6F1-+/+, WBB6F1-W/+ and WBB6F1-W-/+ groups. Groups were distinguished by their coat color. All the mice used in this study were between 6 and 10 wk of age. Food and water were provided ad lib. and, commencing 48 h before killing, the anti-protozoal agent, metronidazole (Poulenc Ltd., Montreal, Canada) was administered in the drinking water (2 g/liter). Experiments designed to assess whether this drug altered NK activity included comparisons of splenic and intestinal NK activity in treated and untreated mice, and the results showed that it did not (unpublished).

Preparation of Effector Cells. Single-cell suspensions from spleen were obtained by mincing and repeated aspiration with pasteur pipettes and filtration through gauze. Lymphocytes from the small intestine epithelium (IEL) and from the lamina propria (LP) were isolated using a modification of previously described procedures (2-8, 16, 17). The small intestines from five mice were removed and flushed of fecal material with 10 ml of saline. The mesentery and adherent connective tissue and fat were dissected from the intestines, and the Peyer’s patches removed. The intestines were opened longitudinally and cut into pieces 1-2-cm long, which were washed in 50 ml of Hanks’ balanced salt solution containing 25 mM Hepes, 5% agammaglobulinemic horse serum (HS), pH 7.4, 300 mosmol/kg (HBSS). To remove epithelial cells and IEL, the tissue was incubated with stirring at 37°C for 15 min in HBSS containing 10^{-4} M EDTA. After supernatant collection, this procedure was repeated once in EDTA-HBSS and once in HBSS. The supernatants were washed twice in HBSS. Histologic examination of the tissue revealed that at this stage, the intestines were devoid of most of the epithelium, although the villus structures and associated LP were largely preserved. To obtain a single-cell suspension from LP, the remaining tissue was cut into 2-5-mm pieces and incubated with stirring at 37°C for 45 min in HBSS containing 25 U/ml of collagenase (840-7018; Gibco Laboratories, Grand Island Biological Co., Grand Island, NY) and 20% HS. After supernatant collection, the collagenase treatment was repeated once. Supernatants from enzymatic digestion were pooled and washed twice. All the cells from the epithelium and the LP were resuspended in 50 ml HBSS and rapidly (10-25 ml/min) filtered through 10-ml syringes containing 300 mg loosely packed nylon wool. This filtration procedure removed a large proportion of the dead cells and debris, and yielded 50-75% of the viable lymphocytes applied to the column. Cells were then washed twice and resuspended in 10 ml of RPMI 1640 supplemented with 10% fetal bovine serum, 10 mM Hepes, and 50/ug/ml gentamycin (Schering Corp. Ltd., Pointe Claire, Canada), pH 7.4, 300 mosmol/kg (growth medium).

At this stage the cell populations usually contained viable epithelial cells, viable lymphocytes, and nonviable epithelial cells (Table I). To obtain enriched lymphocyte populations, a discontinuous density gradient was devised using Percoll (Pharmacia Fine Chemicals, Pharmacia Inc., Uppsala, Sweden), diluted in growth medium (10). Five different concentrations of Percoll were prepared and corrected for osmolarity. The gradient was prepared in 50-ml conical tubes (25339; Corning Glass Works, Corning NY), layering from the bottom: 5 ml 80% (vol/vol) Percoll, 7.5 ml 70% Percoll, 7.5 ml 55% Percoll, 15 ml 40% Percoll and 10 ml 30% Percoll. 2 × 10^8 cells were placed on the top of this gradient, and the tube was centrifuged at room temperature at 600 g for 20 min. Cells from the interface between layers were then collected and washed. Approximately 50% of the viable lymphocytes were recovered. Differential counts of the cell subpopulations were made on smears prepared by cytocentrifugation (Cytospin Centrifuge, Shandon Southern Instruments, Camberley, England) and stained with Diff Quik (Harleco, American Hospital Supply Corp., Gibbstown, NJ).

Cytotoxicity Assay. The YAC-1 clone 19 (18) tissue culture line was obtained through the courtesy of Dr. J. Roder, Queen’s University, Kingston, Ontario and maintained in vitro in growth medium. Tumor cells suspended in 1 ml of growth medium were incubated for 60 min at 37°C with 300 #Ci of Na'~1CrO4 (New England Nuclear, Boston, MA); 10^4 washed cells were then incubated for 6 h at various attacker/target (A/T) cell ratios in 0.7-cm conical bottom wells 76-022-05; Linbro Chemical Co., Hadmen, CT). The percentage of isotope released was calculated from the formula: percent release = (cpm release from cells during incubation –
cpm spontaneous release/[total cpm incorporated × 0.8 - cpm spontaneous release] × 100. Spontaneous release in the 6-h assay was usually between 5 and 12%.

Antisera. Monoclonal anti-Thy-1.2 antibody was purchased from New England Nuclear and used at a final dilution of 1:200 for 10^7 cells/0.5 ml. Monoclonal anti-Lyt-1.1 (lot 7120) and anti-Lyt-2.1 (lot 7301) were purchased from Cedarlane Laboratories (Hornby, Canada). These sera were used at a final dilution of 1:20 and 1:10, respectively, for 10^7 cells/0.5 ml. Anti-NK-1.2 (CE × NZB anti-CBA) alloantiserum (18) was a kind gift from Dr. R. C. Burton, Harvard Medical School, Boston, MA and was used at a final dilution of 1:10 for 10^7 cells/0.5 ml. Anti-asialo GM1 serum produced by Dr. Ko Okumura (19) was kindly donated by Dr. J. Roder and Dr. K. Okumura and used at a final dilution of 1:100 for 10^7 cells/0.5 ml. The antiserum dilutions described above have been previously shown to be effective for removal of the appropriate cell subpopulations (18–20).

Complement (C)-dependent Cytolysis. To kill Thy-1.2, Lyt-1.1, Lyt-2.1, and asialo GM1-positive cells, low toxicity rabbit C (lot 4057; Cedarlane Laboratories) was used at a final dilution of 1:10. To kill NK-1.2-positive cells, rabbit C provided by Dr. R. C. Burton was used at a final dilution of 1:6. Lymphocytes from spleen and intestinal epithelium were incubated at 37°C for 60 min with or without (medium control) the above mentioned antisera in phosphate-buffered saline supplemented with 0.3% bovine serum albumin. Cells were washed once, resuspended in diluted C, and incubated at 37°C for 45 min. Cells were then washed twice and resuspended in growth medium without adjusting the cell concentration, as described by Mattes et al. (21). Viability was assessed by adding fluorescein diacetate and ethidium bromide to the cells, and counting them under UV light (22).

Statistical Analysis. Cytotoxic activity was calculated from 51Cr release assay titration curves by the method of Miller and Dunkley (23) and expressed as lytic units (LU) ± SEM; 1 LU represents the amount of activity required to lyse 20% (LU20) of the target cells during the assay period.

Results

Isolation of Mucosal Lymphoid Cells with NK Activity. We have previously shown that murine IEL possess a high NK cytotoxicity (10). However, the low yield of purified mucosal lymphocytes (≤1×10^6/mouse) obtained by the mechanical isolation procedures used in that study prevented a better characterization of the cells involved. Therefore, we developed a new procedure using EDTA incubations and collagenase digestions for obtaining mouse IEL (5–10×10^6/mouse) and LP lymphocytes (LPL) (~15×10^6/mouse) in high purity. Table I shows the distribution of cells from the intestinal mucosa and fractions obtained by discontinuous density gradient centrifugation after isolation. In the cell populations, before gradient centrifugation, the ratio of viable epithelial cells to viable lymphocytes was from 1:1 to 1:2 for the epithelium, and from 1:3 to 1:4 for the LP. The viable cells represented ~60% of the total cells. As shown in Table I, after the density gradient centrifugations, the majority of viable lymphocytes was found in the 55% and 70% fractions. Interestingly, fraction 55% contained a number of large (>10 μm) gGL, almost twice that of the input population. On the contrary, fraction 70% mainly contained small lymphocytes (~5 μm) with no granules. Thus, the Percoll gradient allowed us to obtain clean populations of lymphocytes enriched or depleted in gGL.

We then assessed IEL and LPL obtained from the gradients for NK activity against YAC-1 tumor cells. Preliminary studies showed that the total IEL and LPL populations possessed marked cytotoxic potential against YAC-1 and that after Percoll fractionation activity was present in the 30% and 55% fractions. However, the input and 30% populations, but not the 55% fraction, expressed high levels of cytotoxicity for NK insensitive targets (MCA/MN fibrosarcoma [10]), and this activity was not
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TABLE I

Distribution of Cells from the Intestinal Mucosa Among Fractions Obtained by Discontinuous Density Gradient Centrifugation

| Source | Fraction | Cells × 10^6 | Percent distribution | Percent granulated lymphocytes in the total lymphoid population |
|--------|----------|-------------|----------------------|---------------------------------------------------------------|
|        |          |             | Viable epithelial cells | Viable lymphocytes | Nonviable cells |
| Epithelium | Input    | 204 ± 33*  | 26 ± 3  | 41 ± 3  | 35 ± 5  | 38 ± 6  |
|          | Percoll 30% | 99 ± 20    | 13 ± 3  | 10 ± 1  | 79 ± 5  | ND*     |
|          | 40%      | 40 ± 16    | 93 ± 7  | 3 ± 3   | 5 ± 2   | ND      |
|          | 55%      | 25 ± 5     | 3 ± 1   | 91 ± 1  | 0       | 64 ± 3  |
|          | 70%      | 6 ± 1      | 0       | 99 ± 1  | 0       | 12 ± 5  |
|          | 80%§     |            |         |         |         |         |
| LP      | Input    | 291 ± 58   | 12 ± 2  | 48 ± 5  | 41 ± 4  | 13 ± 4  |
|          | Percoll 30% | 108 ± 27   | 13 ± 3  | 8 ± 3   | 78 ± 5  | ND      |
|          | 40%      | 64 ± 16    | 46 ± 6  | 23 ± 4  | 26 ± 3  | ND      |
|          | 55%      | 37 ± 12    | 5 ± 2   | 91 ± 2  | 3 ± 1   | 22 ± 4  |
|          | 70%      | 20 ± 9     | 1 ± 1   | 99 ± 1  | 0       | 4 ± 3   |
|          | 80%§     |            |         |         |         |         |

* Mean ± standard error from 10 experiments; five mice per experiment.
† Not done.
§ Contains only bacteria and erythrocytes.

TABLE II

Natural Cytotoxicity Against YAC-1 Tumor Cells of Mouse Lymphocytes from Spleen and Small Intestine

| Source | Percoll fraction | Percent granulated lymphocytes | Percent specific cytotoxicity | LU at 20% level/10^6 cells ± SE |
|--------|-----------------|-------------------------------|-------------------------------|---------------------------------|
|        |                 | 100:1* | 50:1 | 25:1 |                                          |
| Spleen |                 | 30:1  | 20:8 | 10:1 | 2.3 ± 0.2                                  |
| Epithelium | 55      | 32:7  | 19:9 | 12:5 | 2.0 ± 0.3                                  |
|          | 70      | 1:4   | 0:6  | 0:5  | 0.1 ± 0.1                                  |
| LP      | 55      | 19:4  | 14:2 | 9:6  | 1.5 ± 0.2                                  |
|          | 70      | 2:8   | 1:8  | 0:6  | 0:1 ± 0.03                                |

* A/T ratio.
† LU at 20% level/10^6 cells ± SE.
§ Not done.

decreased by freezing and thawing. Therefore, we concluded that the cytotoxic activity of the input and 30% fractions probably reflected nonspecific activity associated with dead cells and debris, and thus we will report results of only the fractions devoid of this (55% and 70% fractions).

Table II shows a representative experiment where statistically significant NK activity was associated with the fractions containing large lymphocytes (fraction 55%). No activity was detected with lymphocytes from fraction 70%. Thus, as previously described for the mouse peripheral blood (13), NK activity correlated with the presence of granulated lymphocytes.
Surface Markers of NK Cells in the Gut Epithelium. To characterize the NK cells in the mouse epithelium, we treated fraction 55% of IEL, which displayed the maximal NK cytotoxicity, with a panel of antisera and C. Splenocytes treated with the same isolation procedures used for intestinal epithelial cells showed the same susceptibility to all antisera plus C as normal splenocytes (data not shown). Table III summarizes the results from the experiments: all antisera killed a proportion of the splenic and IEL cell populations in the presence of C. We then evaluated the effect of the various antisera plus C on NK activity. As shown previously (20), anti-Thy-1.2 plus C reduced splenic NK activity by one-third (Fig. 1). Interestingly, the effect of anti-Thy-1.2 plus C on IEL was much stronger than that on spleen. Treatments with anti-Lyt-1.1 and anti-Lyt-2.1 sera plus C did not affect either the splenic or IEL cytolytic activity (Fig. 2).

Antisera previously reported to be specific for NK cells (18–20) were used with C

| Antiserum     | Percent killed cells ± SE |
|---------------|---------------------------|
|               | Spleen (Percoll 55)       |
| Thy-l,2       | 25.5 ± 2.3                |
| Lyt-1.1       | 28.3 ± 1.2                |
| Lyt-2.1       | 12.1 ± 0.9                |
| Asialo GM1    | 13.5 ± 2.9                |
| NK-1.2*       | 21.4 ± 2.0                |
|               | Epithelium (Percoll 55)   |
| Thy-l,2       | 34.7 ± 4.7                |
| Lyt-1.1       | 21.7 ± 2.3                |
| Lyt-2.1       | 32.6 ± 1.4                |
| Asialo GM1    | 22.4 ± 2.0                |
| NK-1.2*       | 22.1 ± 3.1                |

* The CE × NZB anti-CBA alloantiserum to NK-1,2 kindly provided by Dr. R. C. Burton normally eliminates NK activity at 1:100 to 1:200, but at these dilutions it has little cytotoxic activity on splenocytes. Because our preliminary results showed limited sensitivity of IEL to anti-NK-1.2, we used the reagent at 1:10 to ensure that our results would not merely reflect low levels of NK-1,2 on gut effector cells. At this dilution of the reagent, and using C provided by Dr. Burton, we routinely got levels of cytotoxicity of ~20% (as shown), which probably reflect antibody specificities other than NK-1,2 in the alloantiserum.

![Figure 1](image1.png)

**Fig. 1.** The effect of anti-Thy-1.2 plus C on NK activity by spleen and IEL from Percoll 55%. Percent reduction of L FLU by anti-Thy-1.2 plus C for spleen (○) was 38% (2.4 ± 0.4 LU, C control (△) vs. 1.5 ± 0.1 LU, treated; $P < 0.05$ vs. C), and 75% for IEL (1.5 ± 0.1 LU, C control vs 0.4 ± 0.04 LU, treated; $P < 0.001$ vs. C). (〇), medium alone.
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Fig. 2. The effect of anti-Lyt-1.1 (□) and Lyt-2.1 (○) plus C on NK activity by spleen and IEL from Percoll 55%. Percent reduction of LU₀ by anti-Lyt-1.1 plus C for spleen was −11% (2.4 ± 0.4 LU, C control (△) vs. 2.6 ± 0.2 LU, treated; P < 0.1 vs. C) and 5.5% for IEL (1.5 ± 0.1 LU, C control vs. 1.2 ± 0.1, LU treated; p < 0.1 vs. C). Percent reduction of LU₀ by anti-Lyt-2.1 plus C for spleen was −17% (2.4 ± 0.4 LU, C control vs. 2.8 ± 0.1 LU, treated; P < 0.1 vs. C) and −6% for IEL (1.5 ± 0.1 LU, C control vs. 1.5 ± 0.1 LU, treated P > 0.10 vs. C). (C), medium alone.

Fig. 3. The effect of anti-asialo GM₁ plus C (□) on NK activity by spleen and IEL from Percoll 55%. Percent reduction by anti-asialo GM₁ plus C for spleen was 83% (3.7 ± 0.4 LU, C control (△), vs. 0.5 ± 0.1 LU, treated; P < 0.005 vs. C) and 31% for IEL (2.6 ± 0.4 LU, C control, vs. 1.8 ± 0.3 LU, treated; P < 0.1 vs. C). (○), medium alone.

on spleen and IEL from Percoll gradients. Both anti-asialo GM₁ and anti-NK-1.2 were able to ablate almost the whole NK activity of the spleen (Figs. 3 and 4), but only marginally affected the NK cytotoxicity of IEL. The reduction on IEL by these antisera was not statistically significant (Figs. 3 and 4). Similar results were obtained in all three experiments, which tested in parallel spleen cells and IEL for each antiserum.

**NK Activity of IEL from Beige and W/W<sup>+</sup> Mice.** Using both normal and beige mice (14), Guy-Grand et al. (24) postulated that granulated lymphocytes in the gut are a special class of T lymphocytes capable of acting as progenitors of mucosal mast cells. Because mast cells from mouse peritoneal washings have been recently shown to exert spontaneous cytolytic activity against tumor cells in vitro (25), we decided to investigate the possible relationship between the intestinal NK activity and mast cell
precursors. We used beige mice, known to be deficient in splenic NK activity (14), and W/W<sup>v</sup> mice, known to lack mast cell precursors (15). In agreement with Guy-Grand et al. (24), we found normal numbers of granulated lymphocytes in the mucosa of beige mice. In fact, we observed 46% and 18% granulated lymphocytes in the 55% fractions of Percoll from IEL and LPL, respectively. However, the morphology of the granulated cells was different than normal, because the cells contained only one or
two giant granules (24). When these cells were used in cytotoxicity assays, barely detectable NK activity was observed in IEL and LPL (Fig. 5), and this was similar to the splenic cytotoxic activity.

When we analyzed W/W<sup>v</sup> mice, the percentages of granulated lymphocytes were similar to those observed in the intestines of CBA/J mice, e.g., 66% and 25% gGL were observed in 55% fractions of Percoll from IEL and LPL of W/W<sup>v</sup> mice. Interestingly, when these cells were tested for NK cytotoxicity, low but significant values were obtained for spleen, IEL, and LPL (Fig. 5). To verify whether the lower NK cytotoxicity of W/W<sup>v</sup> mice compared with CBA/J mice was associated with the genetic defects linked to the W/W<sup>v</sup> mutations, littermates from WB/ReJ-W/+ and C57BL/6J-W<sup>v</sup>/+ mice were individually tested for their splenic NK activity. Fig. 6 shows that all the WBB6F1/J mice had a low NK activity regardless of their alleles at the W locus. Thus, the defect in mast cell precursors does not seem to affect the NK activity.

**Discussion**

Using sequential treatments of murine intestinal mucosa with EDTA and collagenase in combination with discontinuous density gradient centrifugation with Percoll, we obtained highly purified populations of lymphocytes from the epithelium and the LP. Because the Percoll gradients allowed us to obtain lymphoid populations enriched or depleted in gGL, we showed that the NK cytotoxic activity in the mouse small intestine (10) was associated with those fractions containing gGL. Even though our results do not prove that gGL are the NK effector cells in the gut, in combination with previous indications that LGL play a central role in the human (11), rat (12), and mouse (13) NK activity, they strongly support the hypothesis that in the gut, spontaneous cytolytic activity against tumor cells is related to granulated lymphocytes.

Apparently, not all LGL are functionally active in the NK cytotoxic assay in vitro, because the proportion of gGL among IEL was usually two- to threefold higher than that among LPL, whereas the NK activity of LPL was usually only slightly lower than that of IEL. Further studies are needed to clarify this point. However, it is of interest that lymphocytes from mouse LP have NK activity, because no spontaneous cytotoxicity was detected by Arnaud-Battandier et al. (7) using guinea pig LPL from the small intestine. Moreover, using human colonic LPL, both the presence and the absence of spontaneous cytotoxicity has been reported (reviewed in 6). Because of these discrepancies with studies in humans, species differences cannot completely explain the discordant results. More likely, technical problems related to the isolation procedures are responsible for the lack of cytotoxicity observed in certain studies. Thus, our positive results in the mouse offer the possibility of performing comparative studies of the different techniques used.

During the course of this study other issues were addressed: (a) is the NK effector cell in the mouse gut identical to the most extensively investigated NK cell in the spleen, and (b) what is the NK lineage?

With a panel of antisera, we observed that lymphocytes with T cell- and NK cell-associated antigens are present in the murine gut mucosa. Only a portion of the IEL from Percoll fraction 55% was killed by Thy-1.2, Lyt-1.1, or Lyt-2.1 monoclonal antiserum plus C (~20–40%). Because it has been reported that the majority (~80–90%) of mouse IEL are Thy-1.2 positive (24), it could be that our isolation procedure
depleted lymphocytes bearing T-associated antigens from fraction 55%. However, variable percentages of T cells, as defined by the E rosette technique, have been reported in IEL from the human colonic mucosa (from 30 to 72%) (26), and in the guinea pig, the proportion of T cells in IEL, as assessed by the E rosette technique, was 48% (26). Thus, the percentage of IEL that bear T cell markers is a controversial issue and will probably be solved only when more standardized isolation procedures are used in all studies (see below).

As previously reported by other authors (20), the splenic NK activity was not affected by anti-Lyt-1.1 and anti-Lyt-2.1 antibodies plus C; identical results were obtained with IEL. Interestingly, anti-Thy-1.2 plus C reduced both the splenic and IEL NK cytotoxicity, but the degree of reduction in the two populations was significantly different. The splenic NK cytotoxicity was reduced by only 30-40%, as shown in other studies (21), whereas the reduction observed with IEL was 75-90%. In contrast, anti-asialo GM1 and anti-NK-1.2, two sera which in the presence of C gave a 70-90% reduction in the splenic NK cytotoxicity, only marginally affected the NK activity of IEL. These results were not caused by different numbers of cells with the corresponding phenotype in gut and spleen, as similar proportions of cells were killed by these two antisera plus C. Accordingly, we suggest that the main IEL effector of NK activity has a phenotype distinct from that of most splenic NK cells. In a previous study (13), we observed that the main effector cells of NK activity in the mouse peripheral blood are LGL. However, similar conclusions could not be drawn for the spleen, where present evidence shows marked heterogeneity in NK activity (27, 28). gGL might be directly derived from LGL in peripheral blood, whose phenotypic characteristics are unknown.

Immunohistochemical studies in tissue sections of murine and human IEL suggest that the majority of the lymphocytes have surface T cell markers (24, 29). In the rat, >50% of IEL have granules (30) and in the rabbit (2), guinea pig (7), and mouse (present study), at least 25% of isolated IEL contained granules. Various studies, including those on the migratory properties of lymphocytes, have suggested that IEL may be derived from T cells originating in Peyer’s patches and migrating to mesenteric lymph nodes, thoracic duct lymph, and the systemic circulation ([24], M. R. McDermott and J. Bienenstock, unpublished observations). The fact that in our hands, NK cells from IEL were Lyt-1−, Lyt-2−, and Thy-1.2+, whereas IEL are said to contain a prominent Lyt-2+ population (31), seems at first sight discordant, but implies that not all IEL have NK activity. Recent studies suggesting major heterogeneity in splenic NK cells may help explain these results, because cells functionally active in NK test systems variably express Lyt antigens and Thy-1 (27). The NK cells we have described in IEL have surface phenotypic characteristics similar to the subset described by these authors (27) and designated NK1. Minato and co-workers (27) speculate that these may be induced to differentiate further to conventional T killer cells. In any event, our studies establish that there is mucosal compartmentalization of a particular subpopulation of NK cells in the intestinal epithelium, and taken together with other evidence for the selective localization in the epithelium of intestinally derived T cells (24), the results support the concept of a unique immune system common to mucosal sites.

Guy-Grand et al. (24) have proposed that the gGL is a mast cell precursor with T cell characteristics. Because Farram and Nelson (25) recently reported that peritoneal
mast cells have natural cytotoxic activity in vitro for certain tumor targets, we investigated the possible relationship between gut NK cells and mast cells. We used two genetically deficient mouse models: the beige, deficient in NK activity (14), and the W/W\textsuperscript{v}, lacking mast cell precursors (15). Extremely low NK activity was found in IEL and LPL from beige mice. W/W\textsuperscript{v} mice had normal numbers of gGL and low but detectable NK activity, suggesting that no lineage relationship exists between NK cells and mast cell precursors. It is important to stress that genetic analysis of the littermates of the WBB6F\textsubscript{i} mice showed no particular NK defect associated with the W/W\textsuperscript{v} alleles. The reason(s) why these results differ from those of Seaman and Talal (32), who showed a depressed NK activity in W/W\textsuperscript{v} relative to littermates, is unclear; perhaps environmental differences affecting one or more subpopulation of NK cells are responsible.

In conclusion, this study has confirmed and extended previous observations (6–8, 10) indicating that granulated lymphocytes from the intestinal mucosa, phenotypically different from the splenic NK cell, are capable of exerting spontaneous cytotoxicity against tumor cells. Lymphocytes with cytoplasmic granules have been described in the epithelium of other anatomical sites such as the mouse lung (10), the rabbit oviductal fimbriae and endocervices (33), and the male reproductive tract of rats and monkeys (34). Thus, the LGL could be a central cell of NK immunosurveillance in several anatomical sites. However, the gGL could have other functions, such as a role in resistance to intestinal infection. Preliminary experiments in mice infected with *Nippostrongylus brasiliensis* or *Giardia muris* show that an increase in splenic NK activity occurs during intestinal infection. Experiments are in progress to assess whether gut mucosal lymphocytes are cytotoxic for intestinal parasites.

**Summary**

Highly purified populations of lymphocytes were obtained from the murine intestinal mucosa using EDTA-collagenase isolation procedures in combination with discontinuous density centrifugation. Intraepithelial lymphocytes (IEL) were separated from lamina propria lymphocytes (LPL) and, within these two populations, fractions enriched or depleted in gut granular lymphocytes (gGL) were obtained. Using these cells in cytotoxic assays, it was shown that both IEL and LPL possess natural killer (NK) activity, and this was associated with gGL. The major effector cells of gut NK activity appeared to be Thy-1.2\textsuperscript{+}, Lyt-1.1\textsuperscript{−}, and Lyt-2.1\textsuperscript{−}. The susceptibility of gut NK cells to anti-Thy-1.2 plus complement (C) was significantly higher than that of splenic NK cells. In contrast, anti-asialo GM\textsubscript{1} and anti-NK-1.2 plus C only slightly affected the gut NK activity. Thus, the phenotype of the gut NK cells appears to be different from the splenic one and provides further evidence for NK heterogeneity and establishes the compartmentalization of one NK subpopulation. Beige mice, deficient in splenic NK activity, also had very low gut NK activity. W/W\textsuperscript{v} mice, which lack mast cell precursors, had normal numbers of gGL and diminished, but still present, gut and splenic NK activity. This deficiency did not segregate with the genes responsible for the basic hemopoietic stem cell defect, and these results argue against a close ontogenetic relationship between IEL, gGL, and intestinal mucosal mast cells. The relevance of these observations to the cell lineage of the effector cell of gut NK activity is discussed.

*Received for publication 22 December 1981 and in revised form 18 March 1982.*
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