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Phenotypic and Functional Separation of Memory and Effector Human CD8\(^+\) T Cells

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Summary

Human CD8\(^+\) memory- and effector-type T cells are poorly defined. We show here that, next to a naïve compartment, two discrete primed subpopulations can be found within the circulating human CD8\(^+\) T cell subset. First, CD45RA\(^-\)CD45R0\(^+\) cells are reminiscent of memory-type T cells in that they express elevated levels of CD95 (Fas) and the integrin family members CD11a, CD18, CD29, CD49d, and CD49e, compared to naïve CD8\(^+\) T cells, and are able to secrete not only interleukin (IL) 2 but also interferon \(\gamma\), tumor necrosis factor \(\alpha\), and IL-4. This subset does not exert cytolytic activity without prior in vitro stimulation but does contain virus-specific cytotoxic T lymphocyte (CTL) precursors. A second primed population is characterized by CD45RA expression with concomitant absence of expression of the costimulatory molecules CD27 and CD28. The CD8\(^+\) CD45RA\(^-\)CD27\(^+\) population contains T cells expressing high levels of CD11a, CD11b, CD18, and CD49d, whereas CD62L (L-selectin) is not expressed. These T cells do not secrete IL-2 or -4 but can produce IFN-\(\gamma\) and TNF-\(\alpha\). In accordance with this finding, cells contained within this subpopulation depend for proliferation on exogenous growth factors such as IL-2 and -15. Interestingly, CD8\(^+\) CD45RA\(^-\)CD27\(^-\) cells parallel effector CTLs, as they abundantly express Fas-ligand mRNA, contain perforin and granzyme B, and have high cytolytic activity without in vitro prestimulation. Based on both phenotypic and functional properties, we conclude that memory- and effector-type T cells can be separated as distinct entities within the human CD8\(^+\) T cell subset.

Recognition of antigen by the immune system evokes a coordinate number of changes in lymphocytes and lymphocyte subsets that allow the system (a) to eliminate or neutralize potential harmful agents and (b) to respond more rapidly and appropriately after renewed antigen encounter, a process referred to as immunological memory. Within the T cell compartment, unprimed (or naïve) T cells (i.e., cells which have not yet encountered antigen), effector T cells (i.e., cells with specialized functions such as cytolysis), and memory T cells can be discerned. Functionally, memory T cells have a number of characteristics that distinguish them from unprimed cells. Not only do they respond efficiently to recall antigens, but, compared to naïve cells, memory T cells also have less stringent requirements for activation and have the potential to secrete a more extensive set of cytokines (1-4). In addition, memory and naïve T cells differ in the expression of several cell surface antigens (5, 6), although it is questioned whether these markers can be looked upon as true memory markers or if they reflect cellular activation (7, 8).

Recent studies by Zimmermann et al. (9) have documented phenotypic distinctions between naïve, effector, and memory CD8\(^+\) T cells in a murine model of lymphocytic choriomeningitis virus infection and an adoptive transfer system with CD8\(^+\) T cells from TCR transgenic mice. Their data confirmed previous reports (10-14) by demonstrating that compared to naïve T cells CTL effectors have strongly downregulated the lymph node homing receptor CD62L (L-selectin) while expression of CD44, CD11a/CD18, CD11b, and CD49d is enhanced. In contrast to effector T cells, memory T cells are heterogeneous with respect to CD62L expression, have no CD11b on their surface, and have an intermediate CD49d expression. Finally, CD45 isoform expression could not be used as a reliable marker for memory-type T cells (9).

In contrast to the murine T cell subsets, description of human CD8\(^+\) naïve, effector, and memory cells is still rather fragmentary. Cytolytic function of CD8\(^+\) cells has been linked to cells with a CD28\(^-\), a CD11b\(^+\), and/or a CD57\(^+\) phenotype (11, 15, 16). Moreover, memory-type
CD8<sup>+</sup> CD45RA<sup>+</sup> CD16<sup>+</sup> T cell subsets that contain enhanced frequencies of antigen-reactive precursor cells (17), have low stringent activation requirements, and have an ability to secrete, apart from IL-2, IFN-γ and TNF-α (18), have been previously described. However, discrimination between naive and memory cells, based solely on CD45 isoform expression, has been proven unreliable, since the CD45RA<sup>+</sup> population contains cells that share a number of phenotypic features with primed cells, such as low CD62L (19) and high CD11a/CD18 (20) expression. Moreover, part of the CD45RA<sup>+</sup> cells have downregulated CD27 (21), a feature considered to result from persistent antigenic stimulation in vivo (22, 23).

In this study we used simultaneous staining with CD45RA and CD27 mAbs to separate functionally distinct subpopulations of human CD8<sup>+</sup> T cells. Our observations demonstrate that, within the human CD8<sup>+</sup> compartment, in accordance with data obtained in experimental animal models, naive, memory- and effector-type cells can be discriminated with coherent phenotypic and functional properties. This type of CD8<sup>+</sup> T cell subset analysis could prove useful in monitoring the immune system in several clinical situations.

Materials and Methods

Reagents. The mAbs CLB-T11.1/1, CLB-T11.2/1, CLB-H1K27 (all CD2), CLB-CD3/3 CLB-CD3/4,1, CLB-CD4/1, CLB-CD8/1, CLB-CD14/1, CLB-Fcγr gran-1 (CD16), CLB-CD19/1, CLB-CD27/1, CLB-CD28, CLB-LFA-1/1 (CD18), CLB-CD49d (VLA-4), and CLB-CD49e (VLA-5), and FITC-conjugated goat anti–mouse Ig, were all produced at the Central Laboratory of the Netherlands Red Cross Blood Transfusion Service (CLB; Amsterdam, The Netherlands). CD95 mAb was purchased from Ortho Diagnostic Systems (Beerse, Belgium). FITC-conjugated–CD62L (Leu-8), –CD57, –CD16, –CD19 (Pharmingen, San Diego, CA), and PE-conjugated CD45RA mAbs to separate functionally distinct subpopulations (see above). Single-stranded cDNA was synthesized from RNA in a 20-μl reaction mixture containing 500 ng oligo(dT) <sub>12-18</sub> and 200 U reverse transcriptase (RT).<sup>1</sup> 1 μl of the reaction mixture was diluted with 11.5 μl of PCR buffer containing 2 mM MgCl<sub>2</sub> and 100 pmol of forward and reverse oligonucleotides. Two PCR reactions were set up for each cDNA, corresponding to Fas ligand (forward: 5'-GGGTCGACCAGGGA-GGTTCAGCTCTTCCACCTAC -3'; reverse: 5'-GCTCATT- GAAACATTCTCGGTCCTGCGTTAACAACGCTTCCATAC -3'; reference 25) and HPRT (hypoxanthine-guanine phosphoribosyltransferase) forward: 5'-TATGGACAGGACTGAAGCTTGTTCTAGCTTCTCCATAC -3'; reverse: 5'-GACCAAAACATGTTCAAACTCCCTGA-3'; reference 26). PCR products were resolved on a 1% agarose gel containing ethidium bromide.

Flow Cytometric Detection of Cytokine Production and Intracellular Staining for Perforin and Granzyme B. Flow cytometric measurement of cytokine production was performed as previously described (27, 28). In brief, 10<sup>6</sup> cells/ml were stimulated for 4 h (IL-4, IFN-γ, and TNF-α) or 8 h (IL-2) with 1 ng/ml PA and 1 μM ionomycin in the presence of the protein-secretion inhibitor monensin (1 μM; all from Sigma Chemical Co., St. Louis, MO). This short-term stimulation did not alter the membrane phenotypic expression of CD45 or CD27 expression. After cell surface staining with PE-conjugated CD45RA and FITC-conjugated granzyme B mAb, provided by Dr. C.E. Hack (CLB), was obtained from Becton Dickinson (San Jose, CA) and IL-12 (donated by Dr. S. Wolf, Genetic Institute, Cambridge, MA).

Cell Preparation and Flow Cytometry. Human PBMCs were isolated from buffy coats of healthy donors by Ficoll-Paque density centrifugation (Pharmacia Biotech AB, Uppsala, Sweden). For phenotypic analysis (Table 1), CD8<sup>+</sup> cells (>95% CD8<sup>+</sup> CD3<sup>-</sup> TCR<sup>-α/β</sup> CD16<sup>-</sup>) were purified by incubating PBMCs with saturating amounts of CD4, CD19, CD16, and CD14 mAbs followed by depletion of positive cells with Dynal beads (Dynal, Oslo, Norway) as previously described (18). Double color immunofluorescence analysis was performed as previously described (23). In brief, purified CD8<sup>+</sup> T cells were incubated with an unlabeled mAb followed by FITC-conjugated goat anti–mouse Ig staining. After blocking free binding sites on the goat anti-mouse conjugate with 10% normal mouse serum, cells were incubated with PE-conjugated CD45RA and biotinylated CD27 mAbs. The latter was then stained with Streptavidin red 670.

For subset purification, CD8<sup>+</sup> T cells were prepared by positive enrichment using the MACS system (Miltenyi Biotec, Bergisch-Gladbach, Germany). In brief, PBMCs (10<sup>7</sup> cells/80 μl incubation buffer; PBS/2% FCS/5 mM EDTA) were stained with CD8 microbeads (Miltenyi Biotec; 20 μl/10<sup>7</sup> positive cells) for 15 min at 4°C. After one washing, cell suspensions were resuspended in incubation buffer (10% cells/400 μl) and enrichment was performed with BS columns (capacity: 10% cells) and the VarioMACS magnet according to manufacturer’s instructions. The resulting CD8<sup>+</sup> T cells were >98% CD8<sup>+</sup> CD3<sup>-</sup> TCR<sup>-α/β</sup> CD16<sup>-</sup> as determined by immunofluorescence analysis with directly labeled mAb. Next, cells were stained with PE-conjugated CD45RA and FITC-conjugated CD27 and sorted into CD45RA<sup>+</sup>CD27<sup>+</sup> (purities: >97%) and CD45RA<sup>−</sup>CD27<sup>−</sup> populations (purities: >97%) on a FACStar®.

Detection of Fas-ligand mRNA. Total RNA was prepared (TRIzol reagent; Life Technologies, Paisley, UK) from 2.5 × 10<sup>6</sup> sorted T cell subsets (see above). Single-stranded cDNA was synthesized from RNA in a 20-μl reaction mixture containing 500 ng oligo(dT)<sub>12-18</sub> and 200 U reverse transcriptase (RT).<sup>1</sup> 1 μl of the reaction mixture was diluted with 11.5 μl of PCR buffer containing 2 mM MgCl<sub>2</sub> and 100 pmol of forward and reverse oligonucleotides. Two PCR reactions were set up for each cDNA, corresponding to Fas ligand (forward: 5'-GGGTCGACGGGA-GGTTCAGCTCTTCCACCTAC -3'; reverse: 5'-GCTCATT-GAAACATTCTCGGTCCTGCGTTAACAACGCTTCCATAC -3'; reference 25) and HPRT (hypoxanthine-guanine phosphoribosyltransferase) (forward: 5'-TATGGACAGGACTGAAGCTTGTTCTAGCTTCTCCATAC -3'; reverse: 5'-GACCAAAACATGTTCAAACTCCCTGA-3'; reference 26). PCR products were resolved on a 1% agarose gel containing ethidium bromide.

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1 Abbreviations used in this paper: B-LCL, B lymphoblastoid cell lines; p, precursors; R.T, reverse transcriptase.
CD27 cells were washed twice with cold PBS and fixated with PBS/4% paraformaldehyde (at 4°C for 5 min). Fixation was followed by permeabilization with PBS/0.1% saponin (Sigma Chemical Co.)/10% human pooled serum (at 4°C for 10 min). For all subsequent incubation and washing steps, PBS/0.1% saponin/0.5% BSA was used. Staining of the cytoplasm with biotinylated cytokine mAbs (IL-2, IL-4, and IFN-γ) at all 5 μg/ml was followed by incubation with Streptavidin red 670 (both at 4°C for 20 min). Biotinylated or FITC-conjugated mouse IgG1 control mAbs (DAKO) were used as negative controls. Data acquisition was performed on a FACScan® and analysis was done using PC-Lysis software.

Intracellular content of perforin and granzyme B was measured in freshly isolated CD8+ T cells without previous stimulation. A permeabilization and staining protocol identical to that described above for the cytokine analysis was used.

Cell Culture and Activation. All cell culture experiments were performed in IMDM supplemented with 10% FCS and antibiotics. Sorted CD8+ T cell subsets (5 × 10⁴ cells/well) were stimulated with a combination of three CD2 mAbs (CLB-T11.1, CLB-T11.2/1, and CLB-HIK27, all 5 μg/ml) in the presence or absence of the following recombinant human cytokines: IL-2 (50 U/ml), IL-4 (10 ng/ml), IL-6 (100 U/ml), IL-10 (100 U/ml), IL-12 (1 ng/ml), IL-15 (10 ng/ml), and IFN-γ (100 U/ml). Proliferation was measured on day 4 by adding 0.2 μCi/well of [3H]thymidine (Amersham, Buckinghamshire, UK) during the last 4 h of culture. All cultures were set up in triplicate.

To measure cytokine secretion, triplicate cultures of sorted cells (2.5 × 10⁴) were stimulated with PMA (100 ng/ml) and ionomycin (1 μM). Supernatant was taken after 24 h for detection of IL-2 and on day 4 for measurement of IFN-γ, TNF-α, and IL-4 content. In our experience, the indicated time points have been proven optimal. IFN-γ, TNF-α, and IL-4 were detected with specific ELISAs (CLB). IL-2 production was measured in a bioassay using IL-2-dependent mouse CTLL cells (29).

Cytotoxicity Assay. CTL activity was determined in a CD3 mAb-mediated cytotoxicity assay as previously described (30). In brief, FcR-bearing P815 target cells were radiolabeled with Na²¹⁵C²⁰O₂ (Amersham) for 30 min at 37°C. Purified CD8+ T cell subsets were incubated with P815 target cells at varying effector/target ratios in the presence or absence of 5 μg/ml CD3 mAb (CLB-CD3/4.1). After a 4 h incubation period at 37°C, the supernatants of triplicate cultures were collected and counted in a gamma counter. Specific cytotoxicity was determined according to the formula: % specific lysis = 100 × [(cpm experimental release - cpm spontaneous release)/(cpm maximal release - cpm spontaneous release)]. The standard error of the mean percentage lysis did not exceed 5%.

Determination of HIV-1-derived Peptide-specific CTL Precursor Frequency. HIV-1-specific CTL precursors (CTLp) were expanded in vitro by Ag-specific stimulation and frequencies of RT derived peptide aa 244-252 (IVLPEKDWS)-specific CTLp were determined as previously described (31, 32). In brief, cryopreserved PBMCs from an HIV-1 seropositive individual (L090; for case report see reference 32) were thawed and sorted into CD8+ CD45RA+ CD27+ and CD8+ CD45RA- CD27+ subsets. Sorted cells were cocultivated with stimulator cells (paraformaldehyde fixed autologous EBV-transformed B lymphoblastoid cell lines [B-LCL] infected with five multiplicity of infection of recombinant vaccinia virus expressing HIV-1, gR T) and irradiated feeder cells (autologous PBMCs; references 31, 32). At days 2 and 9, microcultures were fed with rIL-2 (10 U/ml Proleukin; provided by
Depleted of monocytes, NK cells, and T cells with CD14, CD16, coated with 5 flat-bottomed microtiter plates (Greiner, Nürnberg, Germany) activity was determined by adding cells of the different CD8 subsets as assessed by RT-PCR.

Cord blood

Adult blood

Figure 1. Triple color immunofluorescence analysis of cord blood lymphocytes and PBMCs from an adult donor. Cells were simultaneously stained with either CD4-PerCP or PerCP-labeled CD8 in combination with PE-conjugated CD45RA and FITC-conjugated CD27 mAbs. Cells were gated on forward scatter/side scatter and PerCP-fluorescence. Four distinct populations can be defined within CD8+ cells from adult blood: (1) CD45RA+CD27+ cells, (2) CD45RA-CD27+ cells, (3) CD45RA+CD27- cells, and (4) CD45RA-CD27- cells. Note that there is also a transitional population with dull CD45RA expression. As in the CD4+ subset (23), these cells have an intermediate phenotype for the studied surface markers (data not shown). Functionally, this subset was not analyzed.

Results

Phenotypic heterogeneity of the CD8+ T cell subset. As has been previously noted (21), in contrast to CD4+CD27- T lymphocytes, which are exclusively CD45RA-, within the circulating CD8+ compartment CD27+ T cells can be found within both CD45RA- and CD45RA+ fractions (Fig. 1). These CD8+CD45RA+CD27+ cells are, like CD8+CD45RA-CD27- cells, absent from cord blood (Fig. 1). Therefore, we hypothesize that in response to antigen two separate primed CD8+ populations may develop in humans (a) a population of CD45RA+ CD27+ T cells, and (b) a population of CD45RA-CD27- T cells. This study was performed to analyze whether these phenotypes discriminate between functionally distinct CD8+ subsets.

In a population of healthy adult donors (n = 30), the distribution of the CD8+ T cell subsets defined by simultaneous CD45RA and CD27 staining was as follows (for definition of the subsets see Fig. 1): (1) CD45RA+CD27+ cells 55 ± 17% (range 28–83%); (2) CD45RA-CD27- cells 25 ± 11% (range 9–48%); (3) CD45RA+CD27- cells 13 ± 13% (range 1–50%); and (4) CD45RA-CD27+ cells 4 ± 3% (range 0–15%).

To further define the cell surface phenotype of these subsets, CD8+ T cells were purified from peripheral blood and analyzed in triple color immunofluorescence. As expected on the basis of the observation that these were the only cells present in cord blood, CD45RA+CD27+ cells showed characteristics of unprimed cells since they were CD62L+ and had low CD49d, CD49e, CD29, and CD58 expression. The majority of cells were CD28+ and did not express CD45R0 (Table 1 and data not shown). The CD45RA+ population had features of antigen-primed cells. CD62L was downregulated, and CD49e, CD29, and CD58 were upregulated when compared to the
unprimed population. The main difference between CD27+ and CD27− cells within the CD45RA− population was the expression of CD28. Although the majority of CD27+ cells expressed the antigen, only half of the CD27+ cells were CD28+.

Although CD45R0 was absent from CD45RA+CD27− cells, these T cells had a number of features compatible with a primed phenotype. CD62L was found to be downregulated, whereas CD49d, CD49e, CD29, CD11a/CD18, and CD58 were highly expressed. In comparison to the CD45RA− population, expression levels of CD49d and CD11a/CD18 were even higher on CD45RA+CD27− cells. Remarkably, CD11b and CD57 were present on >50% of the cells and the majority did not express CD28.

Fas (CD95) was not expressed on CD8+CD45RA+CD27+ lymphocytes but could be detected on the majority of CD45RA−CD27− T cells. It should be noted that the antigen density on the latter population was markedly lower than on CD8+CD45RA− T cells (Table 1 and Fig. 2A). Interestingly, Fas ligand mRNA was not present in CD8+CD45RA+CD27+ T cells but was strongly expressed in CD8+CD45RA+CD27− lymphocytes (Fig. 2B, lanes B and C). In contrast, only trace amounts of Fas ligand mRNA were detected in CD8+CD45RA−CD27− cells (Fig. 2B, lane D).

Finally, it should be noted that a CD8+CD27− population expressing low levels of CD45RA can be found in most donors (Fig. 1). Analogous to similar cells within the CD4+ T cell compartment (34), these cells coexpress CD45R0 and were found to have an intermediate membrane phenotype between CD45RA−CD27− and CD45RA−CD27+ T cells (data not shown).

Cytokine Production Capacity of CD8+ T Cell Subsets. In contrast to naïve T cells, which primarily secrete IL-2, the ability to produce a variety of cytokines is a typical feature of primed T cells (35). Cytokine production capacity of CD8+ T cell subsets was measured after stimulation for 4 h with PMA and ionomycin at a single cell level (Fig. 3 and Table 2). CD8+CD45RA+CD27− cells paralleled naïve CD4+ cells in that IL-2 was the main cytokine produced by this subset. In agreement with what would be expected from primed cells, CD8+CD45RA− cells were able to secrete IL-2, IL-4, IFN-γ, and TNF-α. IL-4+ cells were exclusively found within this population with both CD27+ and CD27− cells contributing to the total IL-4 secretion. Also with respect to cytokine expression, the CD45RA+CD27− subset showed characteristics of antigen-experienced cells, in that a high percentage of IFN-γ and TNF-α producers could be detected. Remarkably, in contrast to CD45RA− cells, neither IL-2 nor IL-4 production was found. Since IL-2 production in CD45RA+ cells has been described to reach its maximum later than in CD45RA− cells (36), we also analyzed 8-h stimulated cells. Indeed, the frequency of IL-2 producers increased in the CD45RA+CD27− subset. However, also at this time point, CD45RA+CD27− cells did not produce any IL-2. From Fig. 3 it should be noted that, with respect to the cytokine secretion pattern, cells with a CD27dim expression behave as CD27− T cells in this assay. Furthermore, expression of surface markers on these cells was also comparable to CD27− cells (data not shown).

The results obtained by intracellular cytokine staining were confirmed by measuring the cytokine content of the

Table 2. Intracellular Measurement of Cytokine Production in CD8+ T Cell Subsets

<table>
<thead>
<tr>
<th>Cells</th>
<th>IL-2</th>
<th>IL-2†</th>
<th>IFN-γ</th>
<th>TNF-α</th>
<th>IL-4</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD8+</td>
<td>17±2</td>
<td>16</td>
<td>51±15</td>
<td>48±14</td>
<td>3±1</td>
</tr>
<tr>
<td>CD45RA+CD27+</td>
<td>9±4</td>
<td>27</td>
<td>6±2</td>
<td>12±4</td>
<td>&lt;1</td>
</tr>
<tr>
<td>CD45RA+CD27−</td>
<td>1±1</td>
<td>&lt;1</td>
<td>77±16</td>
<td>69±10</td>
<td>&lt;1</td>
</tr>
<tr>
<td>CD45RA−CD27+</td>
<td>43±17</td>
<td>31</td>
<td>76±9</td>
<td>78±8</td>
<td>6±4</td>
</tr>
<tr>
<td>CD45RA−CD27−</td>
<td>33±14</td>
<td>16</td>
<td>76±12</td>
<td>69±6</td>
<td>9±8</td>
</tr>
</tbody>
</table>

* Cells were stimulated for 4 h, mean percentage of positive cells ± SD, n = 6.
† Cells were stimulated for 8 h, mean percentage of positive cells, n = 2.

Table 3. Cytokine Secretion by CD8+ T Cell Subsets

<table>
<thead>
<tr>
<th>Cells</th>
<th>U/mL</th>
<th>pg/ml</th>
<th>pg/ml</th>
<th>pg/ml</th>
<th>pg/ml</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD45RA+CD27+</td>
<td>100</td>
<td>2,860</td>
<td>370</td>
<td>&lt;1</td>
<td></td>
</tr>
<tr>
<td>CD45RA+CD27−</td>
<td>&lt;10</td>
<td>9,310</td>
<td>2,220</td>
<td>&lt;1</td>
<td></td>
</tr>
<tr>
<td>CD45RA−CD27+</td>
<td>100</td>
<td>11,940</td>
<td>1,660</td>
<td>10</td>
<td></td>
</tr>
</tbody>
</table>

* Sorted cells (2.5 × 10⁴) were stimulated with PMA (100 ng/ml) and ionomycin (1 μM) for 4 d. Cytokine content of the supernatants was tested with a bioassay for IL-2 and ELISAs for IFN-γ, TNF-α, and IL-4.

Figure 3. Intracellular measurement of cytokines in CD8+ T cells. Purified CD8+ cells were stimulated for 4 h (IFN-γ and IL-4) or 8 h (IL-2) with PMA and ionomycin in the presence of the protein secretion inhibitor monensin. After surface staining with CD45 and CD27 mAbs, cells were fixated and permeabilized and intracellular accumulated cytokines were detected with specific mAbs. The histograms show the cytokine staining. Dotted lines indicate the negative controls; the positive cell fraction is painted black; and the dot-plots show the distribution of positive cells (black) among the total cell population (gray).
supernatant after 4 d of culture in the presence of PMA and ionomycin (Table 3). Again, CD45RA⁺CD27⁺ T cells mainly produced IL-2. CD45RA⁻ cells secreted all measured cytokines, whereas CD45RA⁻CD27⁻ cells were only capable of IFN-γ and TNF-α production.

To test if the distinction in cytokine production profiles between the two primed CD8⁺ T cell subpopulations was reflected in their proliferation capacities, three fractions were sorted from purified CD8⁺ T cells, i.e., CD45RA⁺CD27⁺, CD45RA⁻CD27⁺, and CD45RA⁻CD27⁻ cells, and stimulated with a combination of CD2 mAbs supplemented with various cytokines (Fig. 4). The CD45RA⁻CD27⁻ population could not be analyzed because of the low number of these cells in most donors (see above). Culture of the sorted cells with the indicated cytokines alone did not result in proliferation (data not shown). In accordance with a previous report (18) on stringent activation requirements of unprimed CD8⁺ T cells, CD45RA⁺CD27⁻ cells were not able to proliferate in response to CD2 mAbs alone, but addition of IL-2, -15, -16, and -10, and IFN-γ had no effect on proliferation. In contrast, CD2 mAbs alone were sufficient to induce proliferation of CD45RA⁻CD27⁺ cells. IL-2, -15, and -12 further increased proliferation, but IL-4, -6, and -10, and IFN-γ only marginally influenced the CD2-induced response. Proliferation of CD45RA⁻CD27⁻ cells was only observed in the presence of either IL-2 or -15. However, the magnitude of the response was lower than that of the other two populations.

CD45RA⁺CD27⁻, but not CD45RA⁻CD27⁻ T cells, have cytolytic activity without previous in vitro stimulation. Our data thus far have shown that CD45RA⁺CD27⁻ cells possess a number of phenotypic characteristics that have been associated with murine (9) and human (11, 15, 16, 37) cytolytic T cells and that these cells secrete IFN-γ and TNF-α, two lymphokines known to play a role in controlling virus infections (38). Therefore, we addressed the question of whether these cells behave as CTLs in vivo. Purified CD8⁺ cells were sorted into the distinct subsets and their cytotoxic capacity was tested in a CD3 mAb-mediated redirected cytotoxicity assay (Fig. 5). As expected, unprimed CD45RA⁺CD27⁺ cells were not able to lyse the target cells. CD45RA⁻CD27⁺ cells had a low but detectable cytolytic activity at relatively high effector/target ratios. Strikingly, however, the CD45RA⁻CD27⁻ population had potent cytolytic activity.

Fas-Fas ligand interaction is an important mechanism used by CTLs for cell-mediated cytotoxicity (39, 40). Next to this, CTL effectors can kill target cells by releasing a pore-forming protein (perforin) and serine proteases (granzymes), which are stored in intracellular granules, into the vicinity of target cell membranes (41, 42). We analyzed freshly iso-

Figure 4. Proliferative capacity of CD8⁺ T cell subsets. Purified CD8⁺ T cells were sorted into the indicated subsets and stimulated with a combination of three CD2 mAbs in the presence of different cytokines. Proliferation was measured on day 4 by adding [³H]thymidine during the last 4 h of the culture. Results from one out of three experiments are shown.

Figure 5. Cytotoxic capacities of CD8⁺ T cell subsets. Purified CD8⁺ T cells were sorted into the indicated subsets and cytotoxicity was directly analyzed against P815 target cells in the presence of CD3 mAbs in a 4-h ⁵¹Cr-release assay. All effector populations used were unable to lyse P815 cells in the absence of CD3 mAbs (lys <3%), thus excluding N K cell-like activity in this system (data not shown). Results are the mean ± SD of three independent experiments. The distribution of cells in the different subsets was as follows: CD45RA⁺CD27⁺, 18–37%; CD45RA⁻CD27⁺, 7–23%; and CD45RA⁻CD27⁻, 30–38%. (CD8⁺ T cells squares; CD8⁺CD45RA⁺CD27⁺, triangles; CD8⁺CD45RA⁻CD27⁻, plus; CD8⁺CD45RA⁻CD27⁺, circles.)

Figure 6. Expression of mediators of cytotoxicity in CD8⁺ T cell subsets. After surface staining of freshly isolated CD8⁺ cells with CD45RA and CD27 mAbs, cells were fixated and permeabilized and intracellularly present perforin and granzyme B were detected with specific mAbs. The histograms show the staining for the intracellular proteins. Dotted lines indicate the negative controls. The positive cell fraction is painted black. The dot-plots show the distribution of positive cells (black) among the total cell population (gray). Note that the different dot-plot staining is due to the use of either biotinylated CD27 mAb in combination with red 670 or FITC-labeled CD27 mAb.
lated CD8+ cells for the presence of perforin and granzyme B. In accordance with the cytolytic capacities of the different subsets, nearly all cells of the CD45RA+CD27− population contained perforin and granzyme B. Unprimed CD45RA+CD27− cells did not contain these enzymes, whereas low staining could be found within the CD45RA− population (Fig. 6 and Table 4). As already seen in the analysis of cytokine production, cells with dull CD27 expression showed a similar granzyme B and perforin expression pattern to CD27− cells.

The CD45RA−CD27+ Subset Contains Antigen-specific Precursor CTL. CTL effectors exert direct ex vivo cytolytic activity, whereas memory cells need restimulation with antigen to acquire cytotoxic function. Since CD45RA−CD27+ cells, which resemble memory T cells with respect to membrane phenotype and cytokine secretion ability, showed no direct ex vivo cytolytic activity, we studied whether cytotoxic effectors could be generated from this population. To this purpose, PBMCs from an HIV-1 seropositive individual were sorted into CD8+ and CD45RA−CD27+ subsets. CTLs from this individual had been previously found to recognize an HLA-B57 restricted CD8+ T cell compartment, helper and suppressor activity could be ascribed to CD45RA−CD27− and CD45RA+CD27+ T cells, respectively, whereas the CD45RA+CD27− does not appear to directly influence B cell differentiation.

Discussion

In this study, we define discrete subsets (summarized in Table 5) within the CD8+ T cell population based on ex-
CD8 population primarily contains unprimed (naive) cells. The analysis of phenotype and function showed that this, a study by Merkenschlager and Beverley (17) documented an elevated precursor frequency to recall antigens specific for an HIV-1 RT-derived peptide. In line with these studies on the CD8 fraction of all donors. The frequency of these cells in healthy donors is very low (usually <4%) and therefore not accessible for assays that depend on vast quantities of purified cells. However, based on membrane phenotype, cytokine production pattern, and expression of perforin and granzyme B, it is suggested that this population may contain both memory- and effector-type cells.

Based on the finding that the majority of CD8+CD45RA+CD27-T cells lack CD28 expression, it was not surprising to find that this subset shares a number of features that have been previously reported for CD28− lymphocytes (15, 45, 46). Azuma et al. (15) showed that CD8+CD28− T cells are able to lyse FcR-bearing targets in the presence of CD3 mAb. Moreover, Berthou et al. recently reported that most of the perforin containing CD8− lymphocytes lack the CD28 molecule (47). The current data confirm and extend these observations by showing that not only perforin but also granzyme B and Fas ligand are expressed within this T cell subset, which is likely to contribute to the lysis of target cells (39, 40). It is of interest to note that two major costimulatory receptors for T cell growth and cytokine production, CD28 and CD27, are absent from this effector-type T cell. The absence of CD28 implies that this cell type is unable to interact productively (in terms of induction of clonal expansion) with professional antigen-presenting cells. This inability is underscored by the observation that, in marked contrast to both the CD45RA+CD27+ and CD45RA−CD27− subsets, CD45RA+CD27− T cells do not proliferate in response to IL-12, one of the major T cell stimulatory cytokines secreted by activated antigen-presenting cells (48). Finally, absence of CD28 on these cells also precludes costimulatory interactions with other, activated lymphocytes (49). Analysis of the proliferation requirements showed that CD45RA+CD27− T cells responded poorly to most stimuli, which is in agreement with functional studies on the CD8+CD28− T cell subset (15, 46). However, CD45RA+CD27− T cells do respond to IL-2 and to a lesser extent to IL-15, which suggests that in vivo this subset might be influenced by Th-derived signals.

In mice, the expression of CD49d nicely separates naive (low), memory (intermediate), and effector (high expression) T cells, and a study by Christensen et al. has shown that CD49d plays a critical role in efficient homing of effector T cells during virus infections (13). Also, in humans CD49d expression correlates well with the distinct CD8− subsets; CD45RA+CD27− T cells being the ones with the highest expression. Apart from CD49d, β1 integrin interacting with vascular cellular adhesion molecule-1 on activated endothelium and also the high expression of the β2 integrins CD11a and CD11b (Table 1) will probably influ-

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ence the homing properties of these effector-type cells, which may be different from those of naive or memory cells. Indeed, $\text{CD}^8^+\text{CD}45\text{RA}^-\text{CD}27^-$ cells were conspicuously absent from tonsils, whereas $\text{CD}45\text{RA}^-\text{CD}27^-$ cells could readily be demonstrated in this lymphoid tissue (Hammann, D., unpublished data). In line with this observation, it was reported that $\text{CD}11b^+\text{CD}45\text{RBright}$ cells are present in blood, liver, and spleen, but absent from tonsils, lymph node, and thymus (45).

Thus, it would seem that in response to antigen, two types of primed $\text{CD}8^+$ populations may develop. First, an effector-type population with high levels of granzyme, perforin, and Fas ligand expression that exerts cytolytic activity, which has lost the ability for autocrine proliferation but has retained the ability to synthesize IFN-$\gamma$ and TNF-$\alpha$, two cytokines implicated in the neutralization of viruses. Second, a population of memory-type cells, which is not cytolytic without further activation, produces a wide range of cytokines and is able to provide helper activity for B cell differentiation. Kinetic studies in mice showed that effector T cells can be found relatively early after virus infection (8 d), and decline thereafter. In contrast, memory cells become prominent at a later point ($\sim$50 d), and it has been suggested that effector cells seed the memory pool (9, 50).

However, the relation between these subpopulations in humans is unclear at this moment. In major contrast to data obtained in the mouse system, effector-type T cell numbers in humans are relatively stable over time ($\sim$18 months, Rep, M.H.G., unpublished data). In addition, it has been reported that $\text{CD}8^+\text{CD}45\text{RA}^-\text{CD}11\text{abright}$, $\text{CD}11b^+$, and $\text{CD}28^-$ T cells increase with age (45, 51, 52) and, in agreement with this (see Table 5 for phenotypic similarities), we found a similar relationship between age and the percentage of $\text{CD}8^+\text{CD}45\text{RA}^-\text{CD}27^-$ T cells (Rep, M.H.G., unpublished data). The reason for the discrepancy in the kinetics of effector-type T cells between mouse and human is not clear at this moment but it is possible that persistence of antigen, cross-reactive epitopes (53), and/or stimulatory cytokines plays a role (54). It can be imagined that these factors will more readily influence the maintenance of a polyclonal effector T cell population in a natural environment than that of the monoclonal T cell population present in laboratory animals housed under pathogen-free conditions. It is questionable, based on the in vitro experiments, whether effector T cells that are present in the human circulation have a sizable clonogenic potential. Still, it can be envisaged that this type of cell contributes to the inhibition of pathogen spread early in secondary infection through immediate lysis of infected cells.

In agreement with recent data on $\text{CD}28$ subsets by Monteiro et al. (55), we found shortened telomers in $\text{CD}8^+\text{CD}45\text{RA}^-\text{CD}27^-$ compared to $\text{CD}8^+\text{CD}45\text{RA}^-\text{CD}27^-$ in some donors (Hamann, D., K.C. Wolthers, S.A. Otto, P.A. Baars, F.M. Miedema, and R.A.W. van Lier, manuscript submitted for publication). This finding shows that transition to the $\text{CD}45\text{RA}^-\text{CD}27^-$ subset is accompanied by cellular division. Cell cycle analysis showed that, analogous to the $\text{CD}4^+$ population (23, 56), the $\text{CD}8^+$ cells that are in $\text{G}2\text{M}$ can only be found in the minute fraction of $\text{CD}8^+\text{CD}45\text{RBright}$ cells and not in $\text{CD}45\text{RA}^-\text{CD}45\text{R0bright}$ cells (Baars, P.A., unpublished data). This finding implicates that $\text{CD}45\text{RA}^-\text{CD}27^-$ cells could only develop via a $\text{CD}45\text{RA}^-\text{CD}45\text{R0}$ and a subsequent $\text{CD}45\text{RA}^-\text{CD}45\text{R0}$ stage. However, at this moment we cannot exclude that the transition from $\text{CD}27^-$ to $\text{CD}27^+$ takes place in compartments, such as solid tissue, which are not readily available for cell-biological analysis. Finally, similar to what has been described for $\text{CD}28^-$ (57) and $\text{CD}57^+$ (16, 37) T cells, we found a limited usage of $\text{V}$ $\beta$ elements by $\text{CD}8^+\text{CD}45\text{RA}^-\text{CD}27^-$ cells (Hamann, D., K.C. Wolthers, S.A. Otto, P.A. Baars, F.M. Miedema, and R.A.W. van Lier, manuscript submitted for publication). These findings, combined with the data on telomeric repeat length, infer that these circulating effector-type cells in humans are generated through antigen-dependent differentiation and proliferation processes.

Analysis of naive, memory, and CTL effector-type $\text{CD}8^+$ T cells seems relevant not only in patients with chronic viral infections such as CMV, hepatitis B, or HIV-1, but also in patients actively immunized with tumor antigens and after having received an allotransplantation. Patients infected with CMV have been shown to have an increased percentage of $\text{CD}57^+$ cells within the $\text{CD}8^+$ population that contains virus-specific CTLs and can be oligoclonal (16). In HIV-1 infected individuals, an expanded $\text{CD}28^-$ population within the $\text{CD}8^+$ T cell subset has been reported (46). This population has been found to be primarily $\text{CD}57^+$, exerts potent cytolytic activity, and expresses perforin (46, 58). Interestingly, in contrast to healthy donors where $\text{CD}27^-$ and $\text{CD}28^-$ cells form highly overlapping subsets, a large population of $\text{CD}45\text{RA}^-\text{CD}28^-$ cells that still express the $\text{CD}27$ antigen emerges during acute and persistent viral infections (Roos, M.T.L., manuscript submitted for publication), which may point to a sequential downregulation of costimulatory molecules during postthymic T cell differentiation.

Based on the data presented in this paper, we conclude that unidimensional analysis of $\text{CD}8^+$ T cells, using either $\text{CD}45\text{RA}$, $\text{CD}27$, or $\text{CD}28$ mAbs, underestimates the complexity of this subset in humans. Therefore, we propose that combined staining with $\text{CD}27$ and $\text{CD}45\text{RA}$ mAbs provides a tool not only for distinguishing unprimed $\text{CD}45\text{RA}^-\text{CD}27^-$ from primed $\text{CD}45\text{RA}^-$ cells, but also for following the dynamics and expansion of a $\text{CD}45\text{RA}^-$ $\text{CD}27^-$ cytotoxic effector cell population in $\text{CD}8^+$ T cells in different disease states.

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