Immunological memory, defined as faster and more vigorous responses of antigen-primed hosts, is a key characteristic of the adaptive immune system in vertebrates (1). Activation of naive CD4 T cells by antigen induces proliferation and the formation of large numbers of effectors and smaller numbers of memory cells (1–5). However, the lineage relationship of CD4 effector and memory T cells remains controversial (1, 5–8). It has long been debated whether CD4 memory cells can arise from naive precursors either with or without an effector-stage intermediate (9, 10). The corresponding models are termed linear or divergent differentiation models, respectively (1, 5, 6). In the latter case, the effector stage is assumed to be a terminal differentiation stage with the effector cells programmed to die (1, 4–6, 10).

The lineage relationship of CD4 effector and memory T cells has remained difficult to analyze, mostly due to the lack of reliable cell surface markers discriminating between effector and memory stages. Yet, the secretion of effector cytokines constitutes a functional hallmark of effectors (1). Upon antigen activation, naive CD4 cells can differentiate into distinct functional subsets characterized by differential cytokine patterns. Th1 populations produce IFN-γ, whereas Th2 populations secrete IL-4, IL-5, IL-13, and IL-10. Importantly, within short-term in vitro–differentiated or in vivo–generated Th1 and Th2 populations, considerable heterogeneity exists with regard to the production of the respective effector cytokines at the single-cell level (11). This characteristic prevented a clear answer to the question of whether cytokine-secreting effectors can survive the effector stage and develop into...
Previous studies in T cell–deficient mice had shown that antigen and MHC class II molecules were not required to sustain adoptively transferred Th1 and Th2 populations (9, 14–16). However, the recipients’ T cell deficiency could have favored the generation of long-lived cells in unphysiological ways, including the lack of competition with endogenous T cells for survival signals such as cytokines (7, 8). Moreover, the functional properties of memory T cells maintained in the absence of MHC class II interactions were partly impaired upon antigen reencounter (17). Therefore, we have used normal, non-lymphopenic mice as recipients and tested the functional long-lived memory cells, or whether memory cells rather arise from the cytokine-nonsecreting cells within the heterogeneous populations (9, 10). The question is of particular importance because novel immune cell therapies of malignant and infectious diseases in humans focus on the isolation and adoptive transfer of antigen–specific CD4 and CD8 effector T cells identified by antigen-induced secretion of effector cytokines (12). To clarify this issue, here we have purified cytokine-secreting Th1 and Th2 effectors and their nonsecreting counterparts for adoptive transfer into normal mice, followed by the analysis of memory cell formation in vivo (13).
potential of effector-derived Th1 and Th2 memory cells by exposure to cognate antigen in lymphocytic choriomeningitis virus (LCMV) infections. Furthermore, we have analyzed the stability and flexibility of memory Th1 and Th2 cytokine patterns in viral challenge infections and assessed the impact of memory T cell reactivation on LCMV-induced immunopathology. Our results in normal mice show that purified cytokine-secreting Th1 and Th2 effectors can develop into long-lived memory cells with distinct functional properties in vivo, providing direct evidence for the concept of linear memory T cell differentiation.

RESULTS AND DISCUSSION

Generation of memory T cells from purified cytokine-secreting Th1 and Th2 effectors
To investigate whether CD4 effector T cells can survive the effector stage, we purified CD4 effectors secreting effector cytokines for adoptive transfer and analysis of memory cell formation in vivo (Fig. 1 A) (13). Control experiments demonstrated that the reagents used for the purification of cytokine-secreting cells did not affect the survival and maintenance of the cells after adoptive transfer (Fig. S1, available at http://www.jem.org/cgi/content/full/jem.20071855/DC1). To generate T cell populations containing cytokine-producing effectors, naive LCMV-specific CD4⁺Thy1.1⁺ cells were stimulated under Th1- or Th2-differentiating conditions. Within the resulting populations, production of the respective effector cytokines is heterogeneous at the single-cell level (11). Therefore, IFN-γ-secreting Th1 and IL-4⁻ or IL-10⁻ secreting Th2 effectors and their nonsecreting counterparts were identified and sorted by cytometric cytokine secretion assays (Fig. S2) (18). Sorted populations were labeled with CFSE to assess cell divisions after adoptive transfer into previously untreated normal mice. In all recipients, the transferred Thy1.1⁺ cells and/or their progeny were clearly detectable during the next 138 d (Fig. 1, B and C). Sorted cytokine-secreting and -nonsecreting cells showed similar proliferation, half-lives, and persistence over time (Fig. 1 C). Thus, cytokine-secreting Th1 and Th2 effectors gave rise to long-lived memory cells as efficiently as did their respective cytokine-nonsecreting counterparts.

Survival and homeostatic proliferation of CD4 memory cells depend on IL-7R signals (7, 8). At day 2 after transfer, a large proportion of the transferred CD4⁺Thy1.1⁺ cells expressed only low levels of IL-7Rα chain (IL-7Rα), the activation-regulated subunit of the IL-7R complex (Fig. 1 D). By day 5, however, the vast majority of the transferred cells were characterized by high IL-7Rα expression. Notably, IL-7Rα levels correlated positively with the number of divisions that the cells had undergone, which is in line with a role for IL-7 in homeostatic proliferation (7, 8). Cytokine-secreting and -nonsecreting T cells showed a virtually indistinguishable pattern of IL-7Rα expression over time. Thus, IL-7 signals could provide a molecular mechanism for how cytokine-secreting Th1 and Th2 effectors became long-lived and entered the memory pool.

Figure 2. Generation of memory T cells from long-term differentiated Th1 and Th2 effectors. (A) Naive LCMV-specific CD4⁺Thy1.1⁺ cells were differentiated toward Th1 or Th2 for 3 wk with weekly reactivations. Cells were then restimulated with PMA/ionomycin and stained for CD4 and intracellular cytokines. The frequencies of cytokine⁺ cells within CD4⁺ cells are indicated. (B) 3-wk-differentiated Th1 cells (5 x 10⁶) and Th2 cells (2 x 10⁶) were transferred into normal C57BL/6 mice. At the indicated time points, the frequencies of transferred Thy1.1⁺ cells within the peripheral CD4⁺ population were determined by FACS, and the respective t1/2 ± SDs were calculated. Data represent the mean ± SEM of 10 mice per group and time point. Similar results were obtained in two independent experiments.
cells into normal mice and infected the recipients with LCMV. In the course of the infection, virtually all transferred Thy1.1+ cells proliferated extensively (Fig. 3 A). Many of these cells differentiated toward Th1 and produced IFN-γ, TNF-α, and/or IL-2, whereas the numbers of cells producing IL-4, IL-5, IL-13, IL-10, or IL-17 were low (Fig. 3 B). At day 10 of infection, we separated LCMV-specific IFN-γ-secretory effectors and their nonsecreting counterparts from recipients’ splenocytes for further transfer into uninfected normal mice (Fig. 3 C). During the next 101 d, sorted IFN-γ-secreting

Figure 3. Generation of effector memory T cells from purified IFN-γ–secreting effectors primed in an LCMV infection. Naive LCMV-specific CD4+Thy1.1+ cells were labeled with CFSE, and 5 × 10^5 cells were transferred into normal C57BL/6 mice, followed by infection of the recipients with LCMV-Armstrong. (A) At the indicated time points after infection, the frequencies and CFSE dilution of transferred Thy1.1+ cells (red) within the peripheral CD4+ population (blue) were determined. Control recipient mice were left uninfected (naive). (B) On day 7 of infection, splenocytes were stimulated with PMA/ionomycin, and the frequencies of cytokine+ cells within gated CD4+Thy1.1+ donor cells were analyzed by intracellular cytokine staining. (C) On day 10 of infection, LCMV-specific IFN-γ–secreting effectors and their nonsecreting counterparts were identified within splenocytes of the recipients by IFN-γ secretion assay after stimulation with GP64-80 peptide and were sorted by MACS. The frequencies of IFN-γ+ cells within Thy1.1+ donor cells were measured in parallel by intracellular IFN-γ staining (intracellular control). (D) Identical numbers of sorted IFN-γ+ and IFN-γ+/H11002 Thy1.1+ cells as well as IFN-γ−unsorted Thy1.1+ cells (2 × 10^5) were transferred again into naive C57BL/6 mice (n = 3–6 mice per group). At the indicated time points, the frequencies and CD62L expression levels of transferred Thy1.1+ cells within peripheral CD4+ cells were determined. Mean t1/2 ± SDs of CD4+Thy1.1+ cells are indicated. Control mice (no transfer) had received buffer only. (E) Mean frequencies ± SD of CD62Lhi cells within CD4+ Thy1.1+ donor cells were calculated after subtraction of the respective mean Thy1.1 staining background in control mice. (F) 11 d after transfer, LCMV-specific CD8 T cell responses were assessed by tetramer staining and FACS. Memory mice that had been infected with LCMV 21 d earlier and naive C57BL/6 mice served as positive and negative staining controls, respectively.
and nonsecreting cells were maintained with very similar half-lives of 23–28.9 d without indication for a potential survival disadvantage of IFN-γ-secreting effectors (Fig. 3 D). The half-lives of IFN-γ+ and IFN-γ− cells primed in vivo were ≈20 d shorter than the half-lives of sorted IFN-γ+ and IFN-γ− Th1 cells primed in vitro (41.9–48.4 d; Fig. 1 C), possibly reflecting enhanced cell turnover and apoptosis due to more vigorous activation by LCMV in vivo (2, 4, 19, 20). At the time of effector cell sorting, the LCMV-specific CD4 cells displayed an activated CD62Llow, CD44high, CD45RBlow, and IL-7Rαlow phenotype (Fig. S3, available at http://www.jem.org/cgi/content/full/jem.20071855/DC1). Remarkably, upon adoptive transfer, in all groups the vast majority of the transferred T cells stably retained their CD62Llow phenotype over time (Fig. 3, D and E), revealing the preferential generation of effector memory T cells (4). To exclude at best that transfer of sorted CD4 cells had resulted in inadvertent co-transfer of LCMV infectivity, we measured the endogenous LCMV-specific CD8 T cell compartment of the recipients as the most sensitive readout of past LCMV infection. MHC class I tetramer staining failed to reveal LCMV-specific CD8 cells over naive background, showing that viral infection had not been cotransferred (Fig. 3 F). Thus, IFN-γ-secreting effectors primed in vivo in an LCMV infection efficiently gave rise to effector memory T cells maintained in an antigen-independent fashion.

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Figure 4. Unimpaired functional reactivity of Th1 and Th2 effector-derived memory cells upon virus challenge. Naive LCMV-specific CD4+Thy1.1+ cells were differentiated toward Th1 or Th2 for 7 d. Cytokine-secreting and -nonsecreting cells were sorted by cytometric cytokine secretion assays. Unsorted Th1 and Th2 cells and sorted cytokine+ and cytokine− cells (2 × 10⁶) were transferred into normal C57BL/6 mice and rested for 148 d to allow for their differentiation into memory cells. The recipients were then infected with LCMV-WE. (A) At the indicated time points after infection, the frequencies and IL-7Rα expression levels of transferred, Th1-derived CD4+Thy1.1+ memory cells (red) within the peripheral CD4+ population (blue) were determined. (B) Mean frequencies ± SD of transferred Thy1.1+ cells within peripheral CD4+ cells are plotted, and t1/2 ± SDs are indicated. Control mice (no transfer) had received buffer only and were or were not infected with LCMV at day 148. (C) Mean frequencies ± SD of IL-7Rαhigh cells within CD4+Thy1.1+ donor cells were calculated. Control mice that had received unsorted Th1 or Th2 cells were left uninfected and monitored in parallel for IL-7Rα expression levels. Data are representative of two to three independent experiments, each with two to six mice per group and time point (IL-10+ group, one mouse per experiment).
Figure 5. Stability and flexibility of memory Th1 and Th2 cell characteristics without and with virus challenge. (A) Naive LCMV-specific CD4^+Thy1.1^+ cells were differentiated toward Th1 or Th2 for 3 wk before transfer into normal C57BL/6 mice. After 30 d, the frequencies of cytokine^+ cells within gated CD4^+Thy1.1^+ donor cells were analyzed by intracellular cytokine staining upon restimulation of recipients' splenocytes with PMA/ionomycin. (B and C) 30 d after transfer, Th1 and Th2 recipients were infected with LCMV-WE. (B) 7 and (C) 30 d after infection, cytokine production of CD4^+Thy1.1^+ donor cells was measured upon restimulation of splenocytes with PMA/ionomycin. Cytokine analyses of lymph node cells and restimulations with GP{sub}_{64-80}
Antigen responsiveness of Th1 and Th2 effector-derived memory cells upon virus challenge

We tested the antigenic reactivity of effector-derived memory cells by LCMV challenge. When recipients were infected 148 d after adoptive transfer, LCMV-specific memory cells reacted with vigorous proliferation, followed by a protracted contraction phase after virus clearance (Fig. 4, A and B). In recipients of cytokine-secreting and -nonsecreting Th1 and Th2 cells, expansion and subsequent contraction occurred at a very similar rate, with similar kinetics and half-lives, not providing any evidence for defective in vivo expansion of cytokine-secreting effector T cell progeny. Notably, the frequencies of Thy1.1+ memory cells that persisted throughout day 152 after virus challenge ranged from 1.1 to 5.2% (mean 2.3%) of peripheral CD4 cells, which on average exceeded the frequencies before infection by a factor >10 (range 0.2–0.9%; mean 0.2%) (Fig. 4, A and B). Moreover, the kinetics and extent of IL-7Rα regulation were virtually identical for all reactivated memory T cell populations (Fig. 4, A and C). Thus, the usage of IL-7 as a survival factor may provide cytokine-secreting and -nonsecreting T cell progeny with similar capabilities of entering and reentering the memory pool after repeated antigen encounter.

Functional characteristics of Th1 and Th2 effector-derived memory cells upon virus challenge

Little is known about the stability of Th1 and Th2 cytokine patterns in immune responses in vivo. It has been reported that in vitro–differentiated Th1 or Th2 populations largely maintained their respective cytokine profiles when “parked” under neutral conditions in T cell–deficient recipients (9, 14–16). Here, we extend these observations in normal, nonlymphopoenic mice and at the single-cell level. Memory progeny of transferred Th1 cells continued to produce IFN-γ, whereas Th2 memory cells continued to produce IL-4 and other Th2 cytokines upon restimulation after a rest of 30 d in uninfected recipients (Fig. 5 A). However, upon LCMV infection, the Th2 cytokine production profile changed considerably at the population level. Similarly high frequencies of cells of both Th1 and Th2 origin were now found to produce IFN-γ, and this was regardless of whether the memory cells had been generated from sorted IFN-γ+, IL-4+, or IL-10–secreting or –nonsecreting cells (Fig. 5, B and D). Nevertheless, reminiscent of their original differentiation and selectively in cells from Th2 ancestors, a fraction of IFN-γ+ cells coexpressed IL-4 and other Th2 cytokines. Remarkably, these partially differential cytokine patterns, including the substantial number of IL-4/IFN-γ coexpressing cells in the Th2 cell recipients, were stably maintained even in the memory phase 30 and 60 d after LCMV infection (Fig. 5 C and not depicted). Given the extensive proliferation of memory T cells upon LCMV infection and the associated potential for outgrowth of selected subpopulations, the pathways for the generation of such plastic IL-4+IFN-γ+ cells cannot easily be delineated. However, together with previous studies (21–24), our findings raise the possibility that during viral challenge, some cells of Th2 origin still memorize Th2 cytokine expression while acquiring the additional capacity to produce IFN-γ.

Notably, the Th1 or Th2 differentiation history of LCMV-specific memory populations exerted differential effects on T cell–dependent liver disease (25) upon LCMV challenge. All groups of recipient mice with Th1 memory cells developed clear signs of hepatitis, as reflected in elevated serum aminotransferase activities (Fig. 5 E). Yet, none of the groups with Th2–derived memory cells exhibited aminotransferase activities exceeding the levels found in LCMV-infected control mice that had not received LCMV–specific cells, although the reactivated Th2 memory cells expanded vigorously (Fig. 4 B). Virus clearance was not affected in a differential manner by Th1– or Th2–derived memory cells, as LCMV was not detectable in the blood or spleens of any of the recipients at days 7 or 12 after infection, respectively (not depicted). Thus, the partial maintenance of Th2 cell characteristics, such as Th2 cytokine production, may play a role in limiting LCMV–induced immunopathology. Both IL-4 and IL-10 can counteract IFN-γ– and TNF–driven inflammatory processes, possibly by limiting the stimulatory capacity of APCs (26, 27). This may have implications for the activation of endogenous LCMV–specific T cells, including CD8+ CTLs that are most potent in mediating LCMV–triggered liver pathology in mice (25). However, other mechanisms, such as direct cytokine effects, could also be envisaged and will be addressed in future work.

The findings here show that purified cytokine-secreting Th1 and Th2 effectors can efficiently develop into long-lived, functional memory cells, providing direct evidence for the concept of linear memory T cell differentiation. This highlights the similarity of CD4 memory differentiation to analogous processes in CD8 T cell populations (28). Our data, however, do not exclude the possibility of the divergent differentiation model because sorted cytokine–nonsecreting T cells gave rise to memory cells as well. This is compatible with a model in which both linear and divergent differentiation pathways may contribute to the formation of the CD4 memory compartment. Yet, the relative importance of each of the two pathways remains to be investigated. A previous study has reported that IFN-γ+ Th1 effectors tended to be short-lived after adoptive transfer, and memory cells were generated preferentially from IFN-γ– T cells (10). This was interpreted to indicate that the
secretion of IFN-γ demarcates a terminal differentiation stage (10), a view that may appear contradictory to the present data and conclusions. We assume that differences in the experimental systems or procedures could have led to different functional properties of the transferred effector cells, e.g., differential co-production of IL-2 or other survival factors (7, 8, 29) affecting the effectors’ capacity to become long-lived. Using several experimental strategies, our study demonstrates that activation-induced cell death, which can occur in highly activated T cells (19, 20), does not selectively eliminate cytokine-secreting cells.

Importantly, Th1 and Th2 effector-derived memory cells retained functional characteristics reminiscent of their original differentiation. Thus, selected differentiation subsets of antigen-specific effector T cells, adapted to the specific application and selected according to cytokine secretion profile, could be envisaged to optimally target infectious agents or tumor cells and/or to reduce immunopathological side effects of such treatments. Such cells could be provided by immune cell therapy, or they could be induced by appropriately tailored vaccine formulations and adjuvants. Therefore, our finding of efficient generation and maintenance of effector-derived memory T cell populations may open new perspectives in preventive and cell-therapeutic strategies against infectious or malignant diseases.

MATERIALS AND METHODS

Mice. Mice transgenic (tg) for the MHC class II–restricted Smart1 TCR (30) specific for LCMV glycoprotein (GP)64-80 were maintained on the C57BL/6 background under specific pathogen-free conditions and were crossed with Thy1.1+ B6.PL mice. C57BL/6 mice were bred at the Institute of Laboratory Animal Research, University of Zurich, and were used as recipients for adoptive cell transfers at the age of 8–12 wk. All animal experiments were performed in accordance with the Swiss law for animal protection and with permission from the Kantonale Veterinäramt.

Viruses. The LCMV strains WE and Armstrong were grown and propagated as described previously (25), and mice were infected by i.v. injection of 200 PFU in 200 μl.

T cell activation and differentiation. Cell cultures of naive LCMV-TCR Tg CD4+Thy1.1+ T cells and APCs from spleens and lymph nodes were set up in complete RPMI 1640 (cRPMI) as described previously (18). GP64-80 peptide (Neosystem) was used at 0.5 μg/ml. For Th1 differentiation, 3 ng/ml IL-12 and 5 μg/ml anti-IL-4 (11B11; BD Biosciences) were added. For Th2 differentiation, 30 ng/ml IL-4 (Sigma–Aldrich) plus 5 μg/ml anti–IL-12 (C17.8) and 10 μg/ml anti–IFN-γ (R4-6A2; BD Biosciences) were used. Cell cultures were split on days 2 and 4. On day 7, cells were reactivated with fresh Thy1.2-depleted C57BL/6 splenocytes plus GP64-80 peptide, 5 ng/ml IL-2 (R&D Systems), and cytokines and anti-cytokine mAbs as above.

Identification of live IFN-γ-, IL-4-, or IL-10–secreting cells with cytokine secretion assays. Thy1.1+ LCMV-TCR CD4+ cells were activated either with GP64-80 peptide plus Thy1.2-depleted C57BL/6 splenocytes or with 5 ng/ml PMA and 500 ng/ml ionomycin (Sigma-Aldrich). Neutralizing mAbs to IL-4, IL-12, and IFN-γ were added simultaneously at concentrations as above. After 3 h, cells were labeled with an IFN-γ- , IL-4-, or IL-10–specific high-affinity capture matrix (Miltenyi Biotec) as described previously (18). Control samples were either incubated with the respective recombinant cytokine to control the homogeneous labeling of all cells with the cytokine capture matrix (high control), or they were put immediately in ice-cold, instead of 37°C warm, medium to prevent cytokine secretion (low control). The remaining capture-matrix–labeled cells were placed in 37°C warm cRPMI. After 20 (IFN-γ assay) or 30 min (IL-4 and IL-10 assays), matrix–captured cytokine was stained on the cell surfaces with PE-coupled mAb to IFN-γ, IL-4, or IL-10 (Miltenyi Biotec).

Sorting and adoptive transfer of live IFN-γ-, IL-4-, or IL-10–secreting and –nonsecreting cells. After labeling of secreted cytokine with PE-coupled mAb, cells were incubated with anti-PE microbeads (Miltenyi Biotec) and cytokines as above. Samples of unsorted Th1 and Th2 cells were then taken aside. Cytokine– and cytokine–secreting cells were separated by magnetic cell sorting (MACS) as described previously (18). In cases of GP64-80 peptide stimulation, APCs were depleted either by using the CD4 T cell isolation kit (Miltenyi Biotec) or by using a cocktail of biotinylated mAbs to CD8, CD11b, CD11c, CD19, and NK1.1 (all from BD Biosciences), followed by anti-biotin microbeads (Miltenyi Biotec). Sorted purity was controlled on a FACS Calibur (BD Biosciences). Analysis gates were set on live lymphocytes by forward and side scatter and exclusion of propidium iodide–binding particles (0.3 μg/ml; Sigma–Aldrich). Sorted T cells (10^7/ml) were labeled with 5 μM CFSE (Invitrogen) in PBS for 5 min at room temperature. Cells were washed once with cRPMI and then twice with PBS. Identical numbers of Thy1.1+ CD4+ cells from each sorted population were adoptively transferred into previously untreated normal C57BL/6 mice by injection into the tail vein.

FACS analysis of transferred T cells and intracellular cytokines. Transferred T cells were stained with mAbs to Thy1.1 (OX-7), CD4 (RM4-5), CD62L (MEL-14), CD44 (IM7), CD45RB (16A; all from BD Biosciences), and IL-7Rα (A7R34; eBioscience). To prevent unspecific binding of mAbs, all samples were preincubated with 30 μg/ml of blocking anti-FcγRII/III (2.4G2; American Type Culture Collection) and 100 μg/ml of purified rat IgG (Biotrend). Before FACS analysis, erythrocytes were lysed and cells were fixed with FACS lysing solution (BD Biosciences). For analysis of intracellular cytokines, spleen or lymph node cells were stained with IFN-γ, IL-4, and IL-10 mAbs (BD Biosciences) and anti–CD45RB mAb (16A; all from BD Biosciences), and IL-17 mAb (eBioscience). The remaining capture matrix-labeled cells were placed in 37°C warm medium for 6–12 h. Cytokine–secreting cells were stained on the cell surfaces with PE-coupled mAbs to IFN-γ, IL-4, IL-10, IL-17, and IL-12 (R&D Systems). Rat IgG1 (R3-34) and rat IgG2a (R35-95) isotype control mAbs (BD Biosciences) were used at the same concentrations as the respective anti-cytokine mAbs.

Tetramer staining. LCMV-specific CD8 T cells responses to the dominant GP- and nucleoprotein–derived epitopes (GP33 and NP396) were assessed by MHC class I tetramer staining and FACS analysis (30).

Measurement of aminotransferase activity in serum. Aminotransferase activities were determined on a Roche/Hitachi Modular Analytics according to the recommendations of the International Federation of Clinical Chemistry.

Calculation of the half-life. The half-lives (1/2) of memory T cells after transfer and after LCMV infection were obtained by fitting the exponentially decaying function

\[ n(t) = n_0 e^{-\lambda t} \]

to the decaying parts of the measured time series. This function results from the assumption that the number of CD4 T cells of interest declines with a constant net rate λ, so that the cells will disappear eventually (2). \( n(t) \) denotes the cell number at time \( t \), for which data have been measured at distinct time points. The data have been taken into account from time point \( t_0 \), after which the cell number declined exponentially as determined by logarithmic data plotting. The decay rate \( \lambda \) and the initial cell number \( n_0 \) have been chosen.
to obtain the best fit to the data (minimization of the squared differences as implemented in the program package Mathematika). The \( t_{1/2} \) is then

\[
t_{1/2} = \ln 2 / k.
\]

The fits were performed separately for the time series from individual mice. Mean value and SD of \( t_{1/2} \) are shown for the numbers of replicate experiments indicated (first averaging the data of replicate experiments and calculating \( t_{1/2} \) from the averaged data gave very similar results).

**Statistical analysis.** Two groups were compared using an unpaired, two-tailed \( t \) test. More than two groups were compared using one-way ANOVA with Bonferroni’s multiple comparison test. Data from recipients of unsorted and cytokine-sorted T cell subpopulations of either Th1 or Th2 origin were not significantly different (\( P > 0.05 \)) and were combined for statistical analysis.

**Online supplemental material.** Fig. S1 shows that the cytokometric cytokine secretion assay reagents do not affect the survival and long-term maintenance of the labeled cells after adoptive transfer. Fig. S2 shows the sorting of IFN-γ-secreting Th1 and IL-4- or IL-10-secreting Th2 effectors and their nosensecrating counterparts identified by cytokometric cytokine secretion assays. Fig. S3 shows that LCMV-specific CD4 T cells primed by LCMV in vivo acquire an activated CD62Llow, CD44high, CD45RBlow, and IL-7Rαlow phenotype. Figs. S1–S3 are available at http://www.jem.org/cgi/content/full/jem.20071855/DC1.

We thank Edit Horvath for expert technical assistance and Annette Oxenius, Anne Richter, and Mario Assenmacher for helpful discussions and reagents.

This work was supported by the University of Zurich and by grants 3100A0-100779/2, 3100AO-100068/2, and 3100AO-104067/1 of the Swiss National Science Foundation. M. Löhning was a postdoctoral fellow of the Ernst Schering Foundation. M. Swain received support from the Swiss National Science Foundation. J. Hu-Li and M. Assenmacher were supported by the University of Zürich and by grants 3100AO-100068/2, and 3100AO-104067/1 of the Swiss National Science Foundation. M. Swain received support from the Swiss National Science Foundation. J. Hu-Li and M. Assenmacher were supported by the University of Zürich and by grants 3100AO-100068/2, and 3100AO-104067/1 of the Swiss National Science Foundation. M. Swain received support from the Swiss National Science Foundation.

**Submitted: 29 August 2007 Accepted: 13 December 2007**

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