Long-lived memory T cells are able to persist in the host in the absence of antigen; however, the mechanism by which they are maintained is not well understood. Recently, a subset of human T cells, stem cell memory T cells (T_{SCM} cells), was shown to be self-renewing and multipotent, thereby providing a potential reservoir for T cell memory throughout life. However, their in vivo dynamics and homeostasis still remain to be defined due to the lack of suitable animal models. We identified T cells with a T_{SCM} phenotype and stem cell–like properties in nonhuman primates. These cells were the least-differentiated memory subset, were functionally distinct from conventional memory cells, and served as precursors of central memory. Antigen-specific T_{SCM} cells preferentially localized to LNs and were virtually absent from mucosal surfaces. They were generated in the acute phase of viral infection, preferentially survived in comparison with all other memory cells following elimination of antigen, and stably persisted for the long term. Thus, one mechanism for maintenance of long-term T cell memory derives from the unique homeostatic properties of T_{SCM} cells. Vaccination strategies designed to elicit durable cellular immunity should target the generation of T_{SCM} cells.
Superior T memory stem cell persistence supports long-lived T cell memory

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Long-lived memory T cells are able to persist in the host in the absence of antigen; however, the mechanism by which they are maintained is not well understood. Recently, a subset of human T cells, stem cell memory T cells (TSCM cells), was shown to be self-renewing and multipotent, thereby providing a potential reservoir for T cell memory throughout life. However, their in vivo dynamics and homeostasis still remain to be defined due to the lack of suitable animal models. We identified T cells with a TSCM phenotype and stem cell–like properties in nonhuman primates. These cells were the least-differentiated memory subset, were functionally distinct from conventional memory cells, and served as precursors of central memory. Antigen-specific TSCM cells preferentially localized to LNs and were virtually absent from mucosal surfaces. They were generated in the acute phase of viral infection, preferentially survived in comparison with all other memory cells following elimination of antigen, and stably persisted for the long term. Thus, one mechanism for maintenance of long-term T cell memory derives from the unique homeostatic properties of TSCM cells. Vaccination strategies designed to elicit durable cellular immunity should target the generation of TSCM cells.

Introduction

Long-lived memory T cells are able to persist in the host in the absence of antigen (1). In mice, lymphocytic choriomeningitis virus–specific CD8+ T cells are maintained for life after the acute infection (2). Similarly, in humans, vaccinia virus–specific T cells can be found for many decades after vaccination (3). However, it is unclear whether these memory cells are long lived per se, or differentiate regularly from a rarer, long-lived antigen-specific precursor population undergoing slow homeostatic turnover (4).

Dozens of subsets form the memory T cell compartment (5). Conventionally, antigen-experienced T cells have been divided into central memory (TCM) cells and effector memory (TEM) cells, according to their phenotype, function, differentiation history, and anatomical localization (6). Previously, TCM cells as a whole were thought to exhibit stem cell–like behavior, given their capacity to self-renew and to generate more differentiated progeny in response to multiple stimuli (7). However, this concept was recently challenged by the discovery of an earlier stage of memory T cell differentiation in humans, termed stem cell memory (TSCM) (8). TSCM cells are precursors of other memory cells including TEM cells, and display enhanced self-renewal capacity; TSCM cells can also generate multiple subsets of memory cells in vitro, and, despite sharing multiple functional attributes with conventional memory cells, they maintain a largely naive-like phenotype, with a core of expressed genes characteristic of naive cells (8). To date, mouse TSCM cells have been described (9, 10), but those specific for viral or tumor antigens have not been identified, making their relevance in physiology and pathology elusive. To address these questions in a relevant animal model, we attempted to characterize TSCM cells (either as a bulk population or antigen-specific) in healthy nonhuman primates (NHPs) and during the course of SIV infection. The identification of such a population in the NHPs, the most widely used animal model for HIV infection, is directly relevant to the design of an effective HIV vaccine.

Results and Discussion

Human CD8+ TSCM cells display a largely naive-like phenotype, but express high levels of CD95, CXCR3, CD122, and LFA-1 (8, 11). In order to characterize the role of TSCM cells in the generation of T cell memory in vivo, we sought to determine whether a similar subset of cells exists in NHPs. In both healthy rhesus macaques (RMs) and pigtail macaques (PTMs), we identified CD95hi CD8+ T cells in the CD45RA+CCR7+CD27+CD28+IL-7Rα+ naive-like compartment (Figure 1A). Similarly to those in humans, NHP TSCM cells constitute about 2%–3% of circulating CD8+ T cells (Figure 1B). We also identified a CD4+ TSCM subset in PBMCs, with a phenotype and frequency similar to CD8+ TSCM cells (Supplemental Figure 1, A and B; supplemental material available online with this article; doi:10.1172/JCI66327DS1). The NHP model allows a detailed examination of cellular distributions in tissues; we found that CD8+ TSCM cells from healthy RMs are most abundant in LNs, less so in the spleen and bone marrow, and are virtually absent at mucosal surfaces, i.e., the jejunum, the rectum, and the BAL, where only TCM and TEM cells are present (Figure 1C). CD4+ TSCM cells displayed a similar distribution in the body, although less skewed toward the LNs (Supplemental Figure 1C). Thus, TSCM cells have a tropism for secondary lymphoid tissues, with a distribution most similar to naive T (TN) cells.

Conflict of interest: The authors have declared that no conflict of interest exists.

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We next investigated whether NHP TSCM cells have features of memory cells and precede TCM and TEM cells in terms of differentiation. Immunophenotypic analysis of activation and memory markers (8) indicated that NHP CD8+ TSCM cells from healthy RMs are a discrete subset (Figure 2, A and B). Indeed, they are intermediate between TN and TCM cells, according to the expression of proteins that are progressively upregulated (CCR5, IL-18Rα) or downregulated (CD38, CD130) with differentiation (Figure 2B). The human TSCM core phenotypic markers CXCR3, CD122 (Figure 2, A and B), and LFA-1 (Supplemental Figure 2, A and B) were also upregulated in TSCM cells compared with TN cells. Unexpectedly, a relatively large proportion of TSCM cells were proliferating (Ki-67+) in the PBMCs (Figure 2A), but not in the LNs, spleen, or BM (Supplemental Figure 2C). In all sites, TSCM cells were mostly HLA-DR negative. Similar phenotypic data were obtained for CD4+ TSCM cells (Supplemental Figure 1D).

To assess the cytokine production capability of NHP TSCM cells, we stimulated PBMCs from healthy RMs with staphylococcal enterotoxin B (SEB). Following stimulation, all subsets of memory T cells produced IFN-γ, TNF, and IL-2 (Figure 2C). The patterns of cytokine expression show that TSCM cells differ from TCM and TEM cells, with a decreased proportion of cells producing a combination of IL-2 and TNF, but not IFN-γ, and an increased proportion of cells producing IFN-γ and IL-2 or IFN-γ only. Overall, their quality of cytokine production differed from conventional memory T cells (Figure 2C). TSCM cells also display attributes of memory in vivo, as proliferating (Ki-67+) TSCM cells incorporated BrdU following SIVmac239 infection in PTMs similarly to conventional TCM and TEM cells (Supplemental Figure 2D). In contrast, BrdU incorporation by TN cells was negligible.

To further assess whether TSCM cells from healthy RMs constitute discrete, less-differentiated memory cells, we evaluated their multipotency (i.e., to generate other subsets) and self-renewing capability (i.e., to maintain a TSCM phenotype) in response to TCR stimulation in vitro with αCD3/CD28 antibodies (Figure 2D). In the proliferating (CFSE-diluted) fraction, TSCM phenotype cells could only be recovered from sort-purified TN or TSCM cells. In addition, TSCM cells were also able to generate cells with a TCM, TEM, and terminal effector T (TTE) phenotype (Figure 2D). Importantly, TCM cells could not generate TSCM cells, but did generate TCM and TEM cells. Together, these data indicate that TSCM cells constitute a discrete memory subset and support the concept that TSCM cells serve as precursors of other memory cells, according to the relationship TSCM→TCM→TEM→TTE.

To determine the dynamics of antigen-specific T cell subsets during a natural infection in vivo, we enumerated SIV-specific CD8+ T cells in RMs using Mamu-A*01 pMHCI multimers.
presenting SIV-derived Gag CM9 or Tat-TL8 peptides (Supplemental Figure 3A). Antigen-specific T_{SCM} cells were detected within both CM9- and TL8-specific cells at day 21 after infection, thus demonstrating that they are elicited early (Figure 3A). The antigen-specific T_{SCM} cells were not merely “bystander” cells, but displayed evidence of activation (HLA-DR+, CD38bright) and proliferation (Ki-67+) (Supplemental Figure 3B). At day 21 after infection, CM9- and TL8-specific CD8+ T cells were dominated by TEM-like cells, while T_{SCM}, T_{CM}, and T_{TE} cells constituted a small proportion of the total response (Figure 3, A and B).

The biology of the CM9 and TL8 Mamu-A*01–restricted epitopes is highly divergent. Unlike CM9, which is generally maintained intact throughout the chronic phase of infection, the TL8 uniformly undergoes escape mutation within 4–5 weeks after SIVmac239 or SIVmac251 infection in 100% of animals (12–14). Tat-TL8 sequence variants do not stimulate TL8-specific T cells due to their reduced capability to bind the pMamu-A*01 class I molecule (12) or to signal through the TCR (14). Following TL8 escape, there is at least a 10^5-fold decrease in the relative antigen load of TL8 versus CM9. Using this model, it is thus possible to investigate the relative antigen dependence of the different T cell subsets. As expected, viral escape from TL8-directed CTLs resulted in a dramatic decrease in the frequency of total circulating TL8-specific CD8+ T cells between days 21 and 70–120 after infection (Supplemental Figure 3C). Conversely, CM9-specific CD8+ T cells were maintained at high levels throughout the course of the infection (Supplemental Figure 3C). Notably, despite the near-complete elimination of antigen (extremely low levels from residual “archival,” unmutated virus may persist) (12), TL8-specific CD8+ T cells did not disappear, but remained detectable beyond day 335 after infection (Supplemental Figure 3C).

Phenotypic analysis of the TL8-specific population revealed a significant difference in the dynamics of memory subsets in response to viral escape mutation. Indeed, the proportion of TL8-specific T_{EM} cells decreased dramatically in favor of the less differentiated T_{SCM} and T_{CM} cells (Figure 3, A and B). By day 70 after infection, T_{SCM} cells constituted approximately 25% of the total TL8-specific response (Figure 3B). In contrast, memory subset distribution within the CM9-specific CD8+ T cells remained unchanged between days 21 and 70 after infection.
Brief report

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Figure 3

Preferential maintenance of TSCM cells following escape of the cognate antigen. (A) Tetramer+CD8+ T cells (black) superimposed on total CD8+ T cells (gray) at day 21 and day 120 after infection with SIVmac251. Gates first identify CD45RA+CCR7+ cells, then CD28+CD95+ TSCM cells. Numbers in the CD28 versus CD95 plots indicate the percentage of TSCM cells out of total antigen-specific (tetramer+) T cells. (B) Pie charts depicting the proportion of CM9- and TL8-specific CD8+ T cells with different phenotypes (TSCM: CD45RA+CCR7+CD28+CD95+; TCM: CD45RA+CCR7–; TEM: CD45RA–CCR7–; TTE: CD45RA+CCR7–) at day 21 and day 70 after infection (n = 6). **P < 0.01. (C) Fold change in the absolute count of SIV-specific CD8+ T cells between day 21 and day 70 after infection (n = 4). Red bars indicate median values. Data are expressed in log scale. p.i., post infection.
bound annexin V compared with the total memory fraction (there were insufficient T cells for evaluation of TL8-specific TCM and TEM cells individually; Supplemental Figure 6B). Consistent with the lack of chronic antigen exposure, TL8-specific cells were less apoptotic than CM9-specific cells (Supplemental Figure 7B). Notably, genes such as LEF1 (regulating self-renewal), BCL2, and MCLI (both antiapoptotic) were not differentially expressed among “resting” memory subsets, but were specifically upregulated to high levels in “blasting” TSCM cells (Supplemental Figure 7C).

In summary, our data show that NHP TSCM cells are closely related to human TSCM cells, and constitute a discrete memory T cell subset, distinct from TCM cells, on the basis of: (a) surface immunophenotype, (b) localization in the body, (c) cytokine production, and (d) in vivo turnover. Although TCM cells were demonstrated to possess stem cell–like properties in multiple experimental conditions (7), our data suggest that TSCM cells are superior to TCM cells in this regard owing to their superior self-renewing capability and multipotency, their relative antigen dependence in vivo, and their apoptotic refractoriness. Notably, these properties are not unique to TSCM cells, but rather are highly preferentially associated with the TSCM subset. And importantly, the superior persistence of TSCM cells following antigen loss suggests that they are the main precursors of T cell memory in the postantigenic phase. Maintenance of antigen-specific TSCM cells is likely intrinsically programmed. We exclude the possibility that the TSCM pool is maintained by the continuous recruitment of newly generated naive T cells by thymic output (15) in our model, as thymic output is severely impaired in chronic, untreated HIV and SIV infections (16, 17). In addition, we find equivalent maintenance of CM9- and TL8-specific TSCM cells despite a profound difference in antigen availability.

Our data strongly suggest that TSCM cells play a crucial role in supporting long-term cellular immunity in vivo. Future studies aimed at identifying antigen-specific TSCM cells in adoptive transfer models are required to define whether they are uniquely suited for this function. On the basis of these properties, we propose that future vaccination strategies designed to generate durable immunity should target the induction of TSCM cells. Nevertheless, cellular immunotherapy strategies will need to exploit TSCM properties to support the persistence of in vivo–transferred virus- and tumor-specific T cells.

**Methods**

**Animals.** Animals were handled in accordance with the standards of the American Association for the Accreditation of Laboratory Animal Care (AAAACLAC) and meet NIH standards (Guide for the Care and Use of Laboratory Animals). NIH publication no. 85-23. Revised 1985). See Supplemental Methods for animal procedures.

**Antibodies.** Mamu-A*01 tetramers, and flow cytometry. Surface and intracellular analysis of protein expression and cytokine production was performed as described (8, 18). Tetramer staining was conducted at 37°C for 10 minutes. Cells were analyzed and sorted using a modified LSR II and FACSAria II, respectively (BD Biosciences).

**Apoptosis studies.** Fresh PBMCs were cultured in complete RPMI 1640 medium supplemented with cD94d and cD28 antibodies (both at 1 μg/ml; BD Biosciences) at 37°C for 42 hours. Surface phosphatidylserine expression was revealed as described (18).

**Proliferation studies.** Cell proliferation was determined by dilution of CFSE (Life Technologies) as described (8). Labeled cells were stimulated for 6 days with 2 μg/ml CM9 and TL8 peptides in the presence of sorted, CD3+ autologous antigen-presenting cells (1:1 ratio to T cells) or with plate-bound cD93 (10 μg/ml; clone FN18) plus soluble cD28 (1 μg/ml; clone CD28.2) antibodies.

**In vivo BrdU labeling.** Administration of BrdU (Sigma-Aldrich) and analysis of its incorporation were performed as described (18).

**Gene expression analysis.** CD8+ T cell subsets were sorted as depicted in Supplemental Figure 7. See Supplemental Methods for gene expression quantification procedures.

**Statistics.** Analyses were performed using Prism (GraphPad Software Inc.), JMP (SAS Institute Inc.), and SPICE (NIAID, NIH) (19). Nonparametric Wilcoxon rank tests were used to compare distributions. When possible, a paired Student’s t test (2 tailed) was used. In some cases, nonparametric 1-way ANOVA (Kruskal-Wallis test) was used to compare 3 or more groups. Distributions shown with pie charts were compared with SPICE. P values were considered significant when less than 0.05.

**Study approval.** All animal procedures were reviewed and approved by the Institutional Animal Care and Usage Committee of the VRC, the NIAID, the NCI, and Oregon Health and Science University (Portland, Oregon, USA). The anonymous human blood sample (Figure 1A) was obtained from the NIH blood bank under institutional review board exemption.

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