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The sterol regulatory element binding proteins are essential for the metabolic programming of effector T cells and adaptive immunity

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Y.K. designed, did and analyzed most of the experiments and wrote the manuscript; H.E., M.B.H., L.V., K.J.W., J.P.A., B.M.M. E.K. and E.B.W. did experiments and analyzed data; T.F.O. provided materials and intellectual input; and S.J.B. provided overall coordination for the conception, design and supervision of the study and wrote the manuscript (with input from T.F.O., T.G.G., K.R. and D.G.B.).

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Abstract

Newly activated CD8⁺ T cells reprogram their metabolism to meet the extraordinary biosynthetic demands of clonal expansion; however, the signals mediating metabolic reprogramming remain poorly defined. Herein, we demonstrate an essential role for sterol regulatory element binding proteins (SREBPs) in the acquisition of effector cell metabolism. Without SREBP signaling, CD8⁺ T cells are unable to blast, resulting in markedly attenuated clonal expansion during viral infection. Mechanistic studies indicate that SREBPs are essential to meet the heightened lipid requirements of membrane synthesis during blastogenesis. SREBPs are dispensable for homeostatic proliferation, indicating a context-specific requirement for SREBPs in effector responses. These studies provide insights into the molecular signals underlying metabolic reprogramming of CD8⁺ T cells during the transition from quiescence to activation.

Keywords

SREBP; LCMV; lipids; CD8⁺ T cell; metabolism; proliferation

A hallmark of adaptive immunity is the ability of responding T lymphocytes to undergo extensive clonal expansion to protect the host from invading pathogens. To fulfill the requirements of clonal expansion, activated T cells must acquire distinct metabolic programs that meet heightened metabolic demands. Quiescent, naive T cells rely on mitochondrial oxidative phosphorylation (OXPHOS) and catabolic metabolism to fulfill the minimal energetic and biosynthetic requirements associated with quiescence and long-term survival. Upon activation, T lymphocytes rapidly switch their metabolic program to rely predominantly on glycolysis in a process highly analogous to the Warburg effect first described for tumor cells ¹. Additional changes in intermediary metabolism include increased flux through the pentose phosphate pathway (PPP) and an increased reliance on glutaminolysis. The preferential use of glycolysis, pentose-phosphate and glutaminolytic pathways enables rapidly dividing T cells to satisfy the increased metabolic demands during organelle biogenesis and cellular replication ^{2,3}. Accordingly, genetic and pharmacological inhibition of glycolytic, glutaminolytic and lipid biosynthetic pathways attenuate T cell survival, cell cycle progression and effector differentiation ⁴⁻⁹. Despite the importance of metabolic reprogramming for effector cell fate and function, the molecular events linking T cell receptor signaling with metabolic reprogramming remain enigmatic.

A key component in the metabolic reprogramming of effector T lymphocytes is the rapid upregulation of lipid biosynthetic pathways ^{4,8}. Early isotopomer enrichment studies demonstrated that activation of lymphocytes results in the rapid increase of *de novo* cholesterol and fatty acid biosynthesis ⁴. Critically, addition of specific cholesterol derivatives (e.g., oxysterols) to cultures markedly diminished *de novo* lipid biosynthesis and inhibited cell cycle progression in G1, suggesting a link between lipid metabolism and cell cycle progression. Subsequent studies using statins, pharmacologic inhibitors of HMG-CoA reductase (the rate-limiting enzyme in *de novo* cholesterol biosynthesis) also inhibited mitogen-driven lymphocyte expansion ¹⁰. More recently, we and others have established that genetic and pharmacologic perturbations in sterol homeostasis, through the action of the Liver X Receptor (LXR) transcriptional axis, also influence T lymphocyte cell cycle

progression, survival and effector function^{8, 11}. Thus, the regulation of intracellular lipid metabolism is critical for proper lymphocyte growth and function. However, the molecular mechanisms linking mitogenic signaling to the lipid anabolic program of activated lymphocytes remain poorly defined.

The sterol regulatory element binding proteins (SREBP1 and 2) are bHLH-zip transcription factors that have a well-defined role in the regulation of cellular lipid homeostasis¹². In mammals there are two SREBP genes that express three SREBP proteins. SREBP1a and SREBP1c are produced via alternative transcriptional start sites on *Srebf1*, whereas the *Srebf2* gene encodes SREBP2. Canonical SREBP1c signaling preferentially drives expression of fatty acid biosynthesis genes whereas SREBP2 predominately transactivates genes involved in cholesterol biosynthesis, intracellular lipid movement and lipoprotein import. The SREBP1a isoform is able to transactivate both SREBP1c and SREBP2 target genes. In addition to their function in regulating lipid biosynthetic and transport gene expression, SREBPs also transactivate key genes involved in the oxidative PPP and the generation of the co-enzyme NADPH¹³, ensuring sufficient reducing equivalents to meet anabolic demands.

The influence of SREBP signaling on T cell metabolism and function is not well understood. Herein, we use genetic and pharmacologic models to demonstrate that SREBPs are essential for CD8⁺ T cells to undergo metabolic reprogramming in response to mitogenic signaling. Loss-of-SREBP function in CD8⁺ T cells rendered them unable to efficiently blast, resulting in diminished proliferative capacity *in vitro*, and attenuated clonal expansion during viral infection. SREBP activity was required for the upregulation of glycolytic and oxidative metabolism following activation and to maintain lipid levels sufficient to permit rapid membrane biogenesis and cellular growth. The defect in growth and proliferation was traced to a deficiency in cellular cholesterol during blastogenesis which could be rescued by cholesterol supplementation. However, SREBPs were dispensable for homeostatic proliferation, indicating a specific requirement for SREBP signaling to meet the significant anabolic demands supporting effector responses. This study provides mechanistic insights into the metabolic networks ensuring the growth of effector CD8⁺ T cells upon activation, and highlights the importance of SREBP proteins in regulating a cholesterol metabolic checkpoint during blastogenesis.

RESULTS

The TCR induces an mTOR-dependent lipid synthetic program

T cells rapidly upregulate a fatty acid and cholesterol lipid biosynthetic program after stimulation with polyclonal mitogens⁴. To understand the relationship between mitogenic stimulation and induction of the lipogenic program, we activated purified murine T cells with plate bound anti-CD3 alone or in combination with anti-CD28 for 6 hours. Quantitative RT-PCR revealed that anti-CD3 alone was sufficient for induction of cholesterol (*Hmgcr*, *Hmgcs*, *Sqle*) and fatty acid biosynthesis genes (*Acaca*, *Fasn*) (Fig. 1a), and the addition of CD28 costimulation did not provide additional induction of the program. Activation of Pmel TCR transgenic CD8⁺ T cells with increasing doses of agonist gp100 peptide confirmed a requirement for TCR signaling and showed that induction of the lipogenic program occurs at

sub mitogenic doses of antigen (Supplementary Fig. 1a). T cell activation, using the phorbol ester PMA, induced the lipogenic program to levels comparable to that of TCR crosslinking (Fig. 1a). Induction was inhibited by pretreatment of cells with the PKC inhibitor Gö6983 (Supplementary Fig. 1b), suggesting PKC dependence. In contrast, culturing cells with a calcium ionophore (ionomycin) alone did not induce lipogenic genes (Fig. 1a). Taken together, these data indicate that TCR signaling and PKC activation are necessary and sufficient for induction of the lipogenic program in mitogen-stimulated T cells.

Recent studies indicate that loss of Tsc1, a negative regulator of mTOR kinase signaling, resulted in the upregulation of sterol biosynthesis and propanoate metabolism genes in naive CD4⁺ T cells¹⁴. Thus, we asked if induction of lipid biosynthetic programs by mitogens was dependent on mTOR signaling. To address this, purified splenic T cells were pretreated for 30 minutes with 100 nM rapamycin or vehicle and then stimulated up to 6 hours. As expected, activation of T cells robustly induced fatty acid and cholesterol biosynthetic genes, whereas 30 minutes pretreatment of cultures with rapamycin completely inhibited the induction of the lipogenic program (Fig. 1b). Immunoblots on whole cell and nuclear extracts revealed that rapamycin prevented processing of full length SREBP resulting in a failure to accumulate mature SREBP in the nucleus (Supplementary Fig. 1c). Inactivation of the PI3K pathway with LY294002 (Ly294) treatment similarly repressed lipogenic gene expression (Fig. 1b). These data indicate that signaling through the PI3K-mTOR pathway is necessary for the induction of the lipid anabolic gene program of activated lymphocytes.

SREBPs mediate the lipid anabolic program of T cells

To directly address the function of SREBPs in the induction of the lipogenic program by mitogenic signals, purified splenic and lymph node (LN) T cells were retrovirally transduced with a truncated form of SREBP1a or SREBP2 that is constitutively active (designated Δ SREBP1a or Δ SREBP2 herein, Supplementary Fig. 1d,e). mRNA was collected 24 h post infection and expression of the lipid biosynthetic genes were determined by RT-PCR. Both Δ SREBP1a and Δ SREBP2 markedly increased the expression of cholesterol and fatty acid biosynthesis genes when compared to empty vector controls (Fig. 1c), indicating that the lipogenic program of T cells is responsive to SREBP signaling.

In complementary experiments, we asked if SREBPs were required for induction of the lipogenic program by mitogenic signaling. To directly address this, purified mouse spleen and LN T cells were transiently transfected with siRNAs targeting SREBP1 and SREBP2 before stimulation with PMA-ionomycin. After 5 h of stimulation, mRNA was collected and expression of lipid biosynthetic genes assessed. Activation of cells transfected with control siRNA increased expression of genes involved in *de novo* lipid biosynthesis (Fig. 1d). In contrast, siSREBP1 and siSREBP2 transfected cells were unable to upregulate cholesterol synthetic genes (Fig. 1d, Supplementary Fig. 1f). Upregulation of fatty acid biosynthetic genes was inhibited, albeit to a lesser extent. Knockdown of SREBP2 alone was sufficient to inhibit the induction of both cholesterol and fatty acid synthetic genes (Fig. 1d). We were only able to achieve a partial knockdown of SREBP1 (Supplementary Fig. 1f) and correspondingly, we observed a small, but statistically significant effect on fatty acid

synthetic genes (Fig. 1d). However, we were unable to inhibit sterol synthetic genes with this knockdown.

The observation that over-expression of Δ SREBP1a or Δ SREBP2 upregulates both fatty acid and cholesterol biosynthetic genes in activated T cells lead us to hypothesize that SREBP1 and SREBP2 might cooperate, or share occupancy, at the promoters of lipogenic genes. Thus, we performed chromatin immunoprecipitations (ChIP) on SREBP1 and 2 from quiescent and activated T cell lysates. In quiescent cells, SREBP2 was readily detectable at the promoters of *Hmgcr* and *Hmgcs* (Fig. 1e). Activation of T cells resulted in a 10-fold or greater enrichment of SREBP2 at the promoters of *Hmgcr* and *Hmgcs* (Fig. 1e). Clear enrichment of SREBP1 was also detectable at the promoters of *Hmgcr* and *Hmgcs*, canonical SREBP2 target genes. Likewise, we could find both SREBP1 and 2 at the promoters of fatty acid synthetic genes (Fig. 1e). These data suggest that both SREBP1 and 2 may cooperate in driving the lipogenic program of activated T cells. Pre-treatment of cultures with 25-hydroxycholesterol (25-HC, 10 μ M), a well-described sterol inhibitor of SREBP processing markedly reduced enrichment of SREBP1 and 2 at the promoters of lipogenic genes (Fig. 1e), and consequently induction of the lipid synthetic program (Supplementary Fig. 1g). Lastly, pretreating cultures with 100 nM rapamycin also reduced SREBP1 and SREBP2 enrichment to background (Fig. 1e); further indicating that mTOR is required to link TCR signaling with induction of the lipid synthetic program via SREBP.

Deletion of the SCAP inhibits SREBP activity in T cells

Previous studies on SREBP proteins indicate that SREBP1 and 2 can compensate for each other when one protein is deleted^{15, 16}. Moreover, our overexpression and ChIP studies suggested that both SREBP1 and SREBP2 could potentially influence the lipid biosynthetic program in mitogen-stimulated T cells. Thus, we reasoned that loss of both SREBP1 and SREBP2 would be required to fully ablate the lipogenic program of activated T cells. To achieve this, we crossed mice containing floxed alleles of the SREBP chaperone protein SCAP (SREBP Cleavage Activating Protein) with *Cd4*-Cre mice (hereafter referred to as *Scap*^{fl/fl}). SCAP protein is associated with SREBPs in the ER membrane and regulates the processing and subsequent transcriptional activity of both SREBP1 and SREBP2¹⁷. Examination of 5-6 week-old *Scap*^{fl/fl} and littermate control mice revealed no significant differences in the cellularity of thymus, spleen and LN (Fig. 2a). Phenotypic analysis of thymic and peripheral T cell subsets from *Scap*^{fl/fl} and control mice demonstrated no demonstrable differences in the frequency of CD4⁺ and CD8⁺ T cells (Fig. 2b). Nor were there differences in the expression pattern of the activation markers (e.g., CD25 and CD44, Supplementary Fig. 2a), or total numbers of CD4⁺CD25⁺ and CD8⁺CD44⁺ cells observed (Supplementary Fig. 2b). Finally, BrdU pulse chase experiments to follow homeostatic maintenance of peripheral T cells revealed no significant difference in the labeling or decay between control and *Scap*^{fl/fl} mice over a 42 day period (Supplementary Fig. 2c).

Gene expression studies confirmed a near complete deletion of *Scap* in quiescent peripheral *Scap*^{fl/fl} T cells, and we observed a modest change in the expression of lipogenic genes (Fig. 2c). ChIP studies demonstrated that T cell specific deletion of *Scap* modestly reduced the amount of detectable SREBP protein at target gene promoters in quiescent cells (Fig. 2d).

As expected, control cells rapidly upregulated the lipid biosynthetic program (Fig. 2c) and led to increased SREBP1 and SREBP2 at target gene promoters (Fig. 2d). In contrast, the induction of lipogenic genes was markedly attenuated in *Scap*-deficient CD8⁺ T cells (Fig. 2c). Correspondingly, we observed a significant reduction in the amount of both SREBP1 and SREBP2 protein occupying promoters of lipid biosynthesis genes (Fig. 2d). Taken together, our data indicate that the SREBP-dependent lipid biosynthetic program is largely dispensable for quiescent peripheral T cells; however, genetic deletion of *Scap* attenuates the upregulation of the SREBP transcriptome of mitogen-stimulated T cells.

SREBPs influence CD8⁺ T cell growth and proliferation

To determine if loss of SREBP activity would influence T cell blastogenesis and proliferation *in vitro*, purified control and *Scap*^{fl/fl} CD8⁺ T cells were stimulated with anti-CD3 with or without anti-CD28 or IL-2 for 72 h. Control T cells enlarged by 24 h (Fig. 3a and Supplementary Fig. 3a) and diluted CFSE by 72 h after activation (Fig. 3b). In contrast, *Scap*^{fl/fl} CD8⁺ T cells did not undergo significant enlargement (Fig. 3a and Supplementary Fig. 3a), and only modestly proliferated in response to anti-CD3 (Fig. 3b). The addition of exogenous IL-2 and/or anti-CD28, or PMA-ionomycin to cultures provided only a slight or no increase in the proliferation of *Scap*^{fl/fl} T cells (Fig. 3b). Cell cycle analysis indicated that activated *Scap*^{fl/fl} CD8⁺ T cells remained in G0-G1 of cell cycle (Fig. 3c). The addition of CD28 and/or IL-2 to cultures very modestly increased the proportion of cells in S or G2-M phase (Fig. 3c). These data clearly indicate that SREBP signaling is essential for mitogen-driven CD8⁺ T cell blastogenesis, proliferation and survival.

Our cell cycle data indicates that *Scap* deficient T cells remain in G0-G1 phase. To better define whether *Scap*-deficient T cells were capable of moving out of G0 into G1 of cell cycle, we examined the expression of the G1 associated cyclins and restriction point proteins. Immunoblots from control and *Scap*^{fl/fl} CD8⁺ T cells activated for 18h indicated *Scap*-deficient T cells upregulated cyclins D2 and D3 (Fig. 3d). We also observed decreased phosphorylation of the cell cycle checkpoint protein Rb and maintenance of p27^{kip} (Fig. 3d), suggesting that *Scap*-deficient cells were moving into G1 of cell cycle, but were not entering S phase. We also examined if the loss of SREBP activity was resulting in apoptosis. However, kinetic studies indicated no difference in the frequency of cleaved caspase-3-positive cells up to 36 h post-activation (Fig. 3e). Similarly, we observed no difference in the frequency of annexin V-positive *Scap*^{fl/fl} CD8⁺ T cells (data not shown). However, we did observe a significant decrease in the frequency of DAPI-negative cells by 48 h (Supplementary Fig. 3b), suggesting that loss of SREBP activity results in non-apoptotic death late in cultures. Taken together these data indicate that *Scap*-deficient T cells move efficiently into G1 of cell cycle, but are unable to exit before undergoing a caspase-independent cell death.

Loss of SREBP impacts lipid homeostasis of T cells

Next, we asked if genetic ablation of SREBP activity grossly perturbed lipid homeostasis in T cells. To address this, splenic T cells were purified from 5-8 week old *Cd4-Cre* × *Scap*^{fl/fl} or littermate control mice. Total cholesterol and fatty acid (myristate, palmitate and stearate) content were measured by Gas chromatography-Mass spectroscopy (GC-MS) directly *ex*

vivo. *Scap*^{fl/fl} T cells contained an approximate 2-fold decrease in cellular cholesterol compared to littermate control T cells (Fig. 4a). Palmitate was also modestly decreased in quiescent *Scap*-deficient T cells (Fig. 4b), however no significant difference in myristate or stearate was observed in resting cells. Activation of control T cells with anti-CD3-28 for 24 h increased cholesterol, myristate, palmitate and stearate (Fig. 4a,b). In contrast, *Scap*^{fl/fl} T cells were unable to increase their cholesterol, myristate, palmitate and stearate content in response to activation (Fig. 4a,b). Taken together, these data indicate that SREBP signaling is required for defining a lipid homeostatic set point in quiescent T cells, and plays an essential role in meeting the increased lipid biosynthetic demands during blastogenesis.

Loss of SREBP does not perturb proximal TCR signals

Given the importance of cholesterol in plasma membrane fluidity¹⁸, lipid raft formation and TCR signaling¹⁹, we considered the possibility that the diminished proliferative capacity and survival of *Scap*^{fl/fl} CD8⁺ T cells was a function of altered proximal signaling. To address this, T cells were stimulated with anti-CD3 for 60 min and whole cell lysates were probed for phosphorylation of the upstream signaling molecules Zap-70, Lck, Lat and p44/42 MAPK. Regardless of genotype, we observed normal induction of phosphorylation as early as 5 minutes after stimulation, which was maintained out to 60 min (Fig. 4c). We also considered the possibility that we would observe alterations in the PI3K-AKT pathway given the importance of lipids (e.g., myristate) in proper activation of this pathway. However, we did not observe any difference in phosphorylation of AKT and the downstream target S6 in response to activation (Fig. 4d). Moreover, activation with PMA and ionomycin, which obviate the requirement for TCR and co-stimulation, did not provide a significant rescue in the cell cycle progression, viability and proliferative capacity of *Scap*-deficient T cells (Fig. 3b,c). These data suggest that it is unlikely that the inability of *Scap*-deficient T cells to grow is a function of changes in TCR and co-receptor signaling.

SREBPs regulate a gene program required for T cell growth

To better understand how SREBP was influencing CD8⁺ T cell growth and survival, we performed global gene expression studies on quiescent and 6 h stimulated CD8⁺ T cells. A number of genes were either up or downregulated with anti-CD3-CD28 stimulation in control CD8⁺ T cells. Importantly, we observed a very similar pattern of gene expression in activated *Scap*-deficient cells indicating that loss of SREBP activity does not globally perturb T cell activation. Pathway analysis of genes differentially expressed between quiescent control and *Scap*-deficient CD8⁺ T cells revealed a small set of genes involved in cellular survival was downregulated in the absence of SREBP activity (Supplementary Table 1).

Analysis on genes significantly ($P < 0.001$) less-induced in *Scap*-deficient CD8⁺ T cells after activation using DAVID software analysis^{20, 21} revealed significant enrichment in processes associated with sterol biosynthesis, confirming the effect of *Scap*-deficiency in reducing lipid anabolism in the context of T-cell activation (Fig. 5, Table 1, and Supplementary Table 1). Other gene ontology clusters significantly enriched were associated with RNA homeostasis (e.g., RNA processing, ribosome biogenesis and rRNA metabolic processes), nuclear and organelle lumen and nucleolus biogenesis (Table 1). Analysis of predicted

transcription factors mediating differential gene expression between activated control and *Scap*^{fl/fl} CD8⁺ T lymphocytes revealed significant enrichment in SREBP1 and SREBP2, further validating that loss of SCAP protein influences SREBP-mediated lipid anabolism following T cell activation (Table 2). Loss of the transcriptional regulators CREBBP, NFY, and YY1 was also predicted, and since these have well-positioned roles in lipid metabolism, this likely reflect loss of the lipid anabolic program in *Scap*-deficient T cells. Taken together, these data indicate that SREBP signaling specifically regulates a gene program necessary for lipid anabolism and growth of T cells, but does not grossly influence T cell activation.

SREBP is required for metabolic reprogramming

To further interrogate the metabolic state of *Scap*-deficient T cells, we performed cellular bioenergetics analysis on quiescent and mitogen stimulated T cells. No difference in the basal extracellular acidification rate (ECAR) and oxygen consumption rate (OCR) was observed between quiescent control and *Scap*^{fl/fl} CD8⁺ T cells (Fig. 6a). Stimulation of control CD8⁺ T cells with anti-CD3-28 for 24 h resulted in a 10-fold increase in ECAR and a 4-fold increase in OCR. In contrast, *Scap*^{fl/fl} CD8⁺ T cells only modestly increased OCR and ECAR, but were unable achieve the levels of control cells (Fig. 6a).

Next, we directly assessed mitochondrial function in *Scap*-deficient CD8⁺ T cells (ATP production and spare respiratory capacity) by monitoring the OCR in response to sequential treatment with oligomycin (ATPase inhibitor), FCCP (uncoupling agent) and the electron transport chain inhibitors rotenone and myxothiazol (Fig. 6b). Treatment of freshly isolated CD8⁺ T cells with oligomycin, FCCP and rotenone-myxothiazol revealed no difference in basal respiration, ATP production or spare respiratory capacity of quiescent control and *Scap*^{fl/fl} CD8⁺ T cells (Fig. 6c). Activated control CD8⁺ T cells increased basal OCR, mitochondrial ATP production and spare respiratory capacity (Fig. 6c), consistent with significantly heightened mitochondrial function in blasting CD8⁺ T cells. In contrast, activated *Scap*^{fl/fl} CD8⁺ T cells had significant deficiencies in basal respiration, ATP production and spare respiratory capacity (Fig. 6c), suggesting profound respiratory defects. We considered the possibility that the decrease in respiratory capacity was a function of perturbations in mitochondrial mass, however no difference in mitotracker staining was observed between control and *Scap*-deficient CD8⁺ T cells (Fig. 6d). Taken together, these data suggest that SREBP signaling is required for the acquisition of glycolytic metabolism and heightened mitochondrial respiration during CD8⁺ blastogenesis.

Glucose and glutamine are important carbon sources for energetics, anaplerosis (replenishment of TCA cycle intermediates) and macromolecular biosynthesis in activated lymphocytes⁷. Thus, we asked if SREBP signaling was influencing glucose and glutamine utilization in blasting T cells by directly measuring glucose, glutamine and lactate concentrations in culture media 24 hours after activation. Consistent with our ECAR data, we observed a marked decrease in lactate production by activated *Scap*^{fl/fl} CD8⁺ T cells (Fig. 6e). We also observed a corresponding decrease in both glucose and glutamine consumption in activated *Scap*^{fl/fl} CD8⁺ T cells (Fig. 6e). Kinetic analysis of [¹⁸F]-FDG (an ¹⁸F-labeled glucose derivative) uptake by CD8⁺ T cells revealed that both control and *Scap*-

deficient CD8⁺ T cells initially increase their glucose uptake in response to activation (Fig. 6f). However, *Scap*-deficient CD8⁺ T cells were unable to increase glucose uptake, resulting in an approximate 50% reduction in [¹⁸F]-FDG signals at 18 hours after activation.

RT-PCR analysis revealed that *Scap*^{fl/fl} CD8⁺ T cells have significantly lower expression of glycolysis genes (Supplementary Fig. 4a). Both Myc and HIF1 α have important roles in the upregulation of glycolytic program²², thus we considered the possibility that *Scap*-deficient T cells might have perturbations in these pathways. However, immunoblots indicated normal upregulation of Myc, and increased stabilization of HIF1 α in activated *Scap*-deficient CD8⁺ T cells (Supplementary Fig. 4b). Thus, the inability of *Scap*^{fl/fl} CD8⁺ T cells to acquire a glycolytic program during activation is likely not a direct function of altered Myc or HIF1 α signaling.

A logical prediction of our metabolic studies is that blasting *Scap*-deficient T cells would have altered bioenergetics. To directly address this, we determined intracellular ATP concentrations in quiescent and anti-CD3-28 activated CD8⁺ T cells. No difference in ATP concentration was observed between control and *Scap*^{fl/fl} CD8⁺ T cells directly *ex vivo* (Fig. 6g). Activation of control CD8⁺ T cells modestly increased ATP levels by 6 h, and resulted in a 4-fold increase in cellular ATP by 24 h (Fig. 6g). In contrast, *Scap*^{fl/fl} CD8⁺ T cells did not increase their cellular ATP concentration at 6h and increased ATP by 2-fold at 24 h post stimulation (Fig. 6g). We considered the possibility that perturbations in ATP levels would activate the AMP-activated protein kinase (AMPK) pathway in *Scap*-deficient CD8⁺ T cells. Consistent with earlier studies²³, activation of control T cells resulted in phosphorylation of AMPK and the AMPK target acetyl-CoA carboxylase (ACC). No difference in p-AMPK and p-ACC from *Scap*^{fl/fl} T cells was noted (Supplementary Fig. 4c), suggesting that the difference in glycolysis and cellular growth are not a function of exaggerated AMPK signaling. Taken together, these data support the notion that SREBP signaling influences cellular bioenergetics during blastogenesis independent of the Myc, HIF1 α and AMPK pathways.

Cholesterol restores growth of *Scap*-deficient T cell

Our gene expression data indicates that loss of *Scap* primarily influences lipid homeostasis, in particular flux through the mevalonate pathway. In addition to cholesterol synthesis, the mevalonate pathway is responsible for the generation of non-steroidal lipid modification to proteins, such as prenylation (e.g., Ras)^{8, 24}. Thus, we asked if addition of excess mevalonic acid to cultures would increase flux into isoprenoid synthesis, and rescue cellular growth to *Scap*-deficient T cells. However, the addition of mevalonic acid (50-100 μ M) did not impact survival or blastogenesis (Supplementary Fig. 5a), suggesting that a defect in the prenylation of proteins could not mechanistically explain the phenotype of *Scap*^{fl/fl} CD8⁺ T cells.

These data led us instead to posit that cholesterol was limiting in *Scap*^{fl/fl} CD8⁺ T cells. To directly test this prediction, control and *Scap*-deficient T cells were activated in complete media supplemented with 5 μ g/ml of cholesterol conjugated to Methyl- β -cyclodextrin (MBCD-Chol). Analysis of cultures up to 48 h revealed that addition of cholesterol to control CD8⁺ cells provided a modest increase in cellular survival, with no influence on size (Fig. 7a, Supplementary Fig. 5b). In contrast, *Scap*-deficient T cells doubled their survival at

48 h from 21 % to 45%, approaching that of control T cells without supplemental cholesterol (54%). Furthermore, supplementation with cholesterol restored the cellular enlargement of *Scap*^{fl/fl} CD8⁺ T cells following activation with anti-CD3-28 (Fig. 7a,b). Analysis of cell cycle 48 h after stimulation indicated that addition of cholesterol markedly increased the frequency of *Scap*^{fl/fl} CD8⁺ T cells in S and G2-M (Fig. 7c). Likewise, CFSE analysis indicated that *Scap*-deficient T cells were now able to undergo robust proliferation in response to mitogenic stimulation (Fig. 7d).

Cellular cholesterol levels have been implicated in regulating phospholipid synthesis. The observation that replenishing cholesterol was sufficient to significantly restore the growth of *Scap*-deficient T cells led us to ask if membrane biogenesis was diminished in the absence of SREBP activity. Membrane biosynthesis occurs in the endoplasmic reticulum (ER) before shuttling to other intracellular organelles or the plasma membrane. Thus, we assessed total ER content by flow cytometry in quiescent and newly activated cells using ER tracker. Quiescent *Scap*^{fl/fl} CD8⁺ T cells had decreased ER content when compared to their control counterparts (Fig. 7e). Activation of control cells led to the addition of ER (Fig. 7e). In contrast, *Scap*^{fl/fl} CD8⁺ T cells were unable to expand the ER during blastogenesis (Fig. 7e). However, the addition of exogenous cholesterol restored ER growth (Fig. 7f). These data are consistent with previous work implicating SREBPs in membrane biogenesis²⁵, and support the notion that intracellular cholesterol serves as a metabolic checkpoint in cell cycle progression via regulation of ER membrane biogenesis.

SREBPs intrinsically regulate virus-specific immunity

To determine if SREBP signaling was also required for efficient T cell clonal expansion *in vivo*, *Scap*^{fl/fl} and control mice were challenged with LCMV Armstrong (Arm). LCMV-Arm infection of wild type mice generates a robust virus-specific T cell response leading to rapid control of virus infection within 10-12 days²⁶. On day 8 following infection, splenocytes were counted and stained with LCMV-specific D^bGP33 tetramers to determine the level of virus-specific CD8⁺ T cell. *Scap*^{fl/fl} mice exhibited a significant decrease in the frequency and total number of D^bGP33-specific CD8⁺ T cells (Fig. 8a-c). A similar decrease was observed in the D^bNP396-specific CD8⁺ T cell, indicating a globally diminished antiviral T cell response (Fig. 8c). Intracellular interferon- γ (IFN- γ) and tumor necrosis factor (TNF) staining in response to GP33 peptide challenge *ex vivo* confirmed a decrease in the frequency of responding CD8⁺ T cells (Fig. 8d,e). The few LCMV-specific CD8⁺ T cells that were generated in *Scap*-deficient mice, displayed decreased expression of IFN- γ and TNF (Fig. 8d), suggesting a potential functional defect in the absence of SREBP activity.

Finally, our data indicates an essential requirement for SREBPs in mitogen-driven proliferation *in vitro* and clonal expansion of viral-specific CD8⁺ effector cells *in vivo*. However, we did not observe significant perturbations in the frequency or absolute numbers of peripheral T cells from naive *Cd4-Cre-Scap*^{fl/fl} mice (Fig. 2). These data led us to posit that SREBP activity might not be required under homeostatic conditions. To initially address this, we examined the homeostatic proliferation of purified, CFSE-labeled CD8⁺ T cells adoptively transferred in to Thy1.1⁺RAG2-deficient hosts. On day 6 following adoptive transfer, CFSE dilution of Thy1.2⁺ T cells from LN and spleen was assessed. Consistent

with our hypothesis, we observed no difference in the proliferative patterns of *Scap*-deficient and control T cells (Fig. 8f). Similar proliferative results were seen in a sub-lethal irradiation model of homeostatic proliferation²⁷ (Fig. 8f). RT-PCR revealed that homeostatically proliferating cells did not upregulate the SREBP driven cholesterol synthetic genes (Fig. 8g), although we did observe a very modest upregulation of *Fasn*. Thus we posited that a major difference between T cells undergoing homeostatic proliferation versus conventional antigen-driven proliferation might be cellular growth. Indeed, analysis of cell size indicated that homeostatically proliferating OT-I cells remained small despite undergoing significant proliferation whereas OVA immunized OT-I cells underwent conventional blastogenesis (Fig. 8h). Taken together, these data reveal a context specific requirement for SREBP signaling during conventional antigen-driven responses to support cellular enlargement and effector function.

DISCUSSION

The molecular events guiding the metabolic reprogramming of T cells during activation remain poorly understood. Our studies convincingly demonstrate that SREBPs are essential to coordinate TCR signaling with a lipid anabolic program requisite for rapid membrane biosynthesis and cellular growth. In the absence of SREBP signaling, newly activated CD8⁺ T cells move into G1 of the cell cycle but are unable to grow, resulting in decreased proliferative capacity and markedly attenuated anti-viral immune responses. Strikingly, this defect in growth and proliferation was traced to an insufficiency in cholesterol. These data support a model where an intracellular cholesterol pool, likely in the endoplasmic reticulum – the site of lipid biosynthesis – serves as a metabolic signal conveying the fitness of a blasting T cells. Consequently, the SREBP gene program facilitates the transition of mitogen-stimulated T cells through a critical metabolic checkpoint during blastogenesis. These signals ensure coordination of cellular metabolism with cell cycle. Simply put, without sufficient cholesterol, effector T cells are unable to grow in response to conventional antigen stimulation.

One quite surprising aspect of our study is that SREBP activity is dispensable for homeostatic proliferation of CD8⁺ T cells. SREBPs have importance in regulating the proliferative capacity and survival of cancer cells^{28, 29}, albeit through poorly described and disparate mechanisms. The influence of SREBPs on primary cell proliferation is even less well understood. Our studies on conventional antigen-driven proliferation and anti-viral immunity clearly support the concept that SREBPs influence rapid cellular growth. Nevertheless, the observation that *Scap*-deficient T cells have no observable defects in homeostatic proliferation is the first clear evidence that there are differential requirements for SREBP signaling during proliferation. As such, the emerging notion that engagement of the SREBP pathway is universally required for cellular proliferation and survival is likely incorrect. Alternatively, we would propose that SREBP signaling is necessary under circumstances where cells require the rapid addition of biomass, rather than proliferation per se.

An interrelated and important concept centers on the observations that effector and memory CD8⁺ T cells employ distinct metabolic programs. Generation of effector CD8⁺ T cells

appears to be favored by glycolytic and anabolic programs, whereas memory T cell generation is enhanced under heightened OXPHOS and catabolism^{2, 30}. Our metabolic studies indicate that SREBP signaling is required for a CD8⁺ T cell to fully acquire a glycolytic phenotype. One intriguing possibility is that inhibition of SREBP in responding T cells could preferentially enrich for memory at the expense of effector cell generation. Recent studies have indicated that moderate inhibition of mTOR signaling with low dose rapamycin favors the generation of memory CD8⁺ T cells^{31, 32}. Other studies indicate that mTOR signaling controls cellular metabolism through the HIF and SREBP transcription factors³³, and the data presented here clearly indicate that SREBP in CD8⁺ T cells is rapamycin-sensitive during T cell activation. A logical prediction of these studies is that low dose rapamycin is favoring the development of CD8⁺ memory cells through its effects on SREBP activity. Future studies will be required to rigorously test this prediction. Nevertheless, our data raise the possibility that generation of memory and effector cells could be uncoupled via manipulation of sterol metabolism.

In conclusion, these studies delineate a role for SREBP in controlling conventional antigen-driven clonal expansion, but not homeostatic proliferation. These observations could have significant implications for developing novel therapeutic strategies targeting this pathway to control unwanted effector T cell proliferation, without perturbing homeostasis or development.

Methods

Mice

Scap^{flox/flox}¹⁷ and C57/BL6 mice were purchased from the Jackson Laboratory. *Cd4*-Cre mice were purchased from Taconic. *Scap*^{flox/flox} mice were crossed with *Cd4*-Cre mice to generate *Cd4*-Cre/*Scap*^{flox/flox} mice. *Cd4*-Cre/*Scap*^{+/+} mice were used as littermate control for all experiments using *Cd4*-Cre/*Scap*^{flox/flox} mice. All mice were maintained in pathogen free facilities of the University of California, Los Angeles. All experiments on mice and tissues collected from mice were performed in strict accordance with University of California, Los Angeles policy on the humane and ethical treatment of animals.

Cell culture and reagents

Cells were cultured in IMDM supplemented with 10% heat inactivated FBS, 100 U/ml penicillin, 100 µg/ml streptomycin, 50 uM 2-ME. Total T cells or CD8⁺ T cells were purified using negative enrichment kit (STEMCELL Technologies). T cells were cultured in 96-well plates coated with 5 µg/ml of anti-CD3 (2C11;Bio X Cell) alone or in combination with 2 µg/ml of soluble anti-CD28 (37.51;Bio X Cell) or 100 IU/ml of human IL-2. Alternatively, cells were cultured with 50 ng/ml of PMA (EMD Chemicals) alone or in combination with 500 ng/ml of ionomycin (Sigma-Aldrich). For proximal TCR signaling analysis, splenocytes were cultured with 1 µg/ml of soluble anti-CD3. In some experiments, cells were cultured with following inhibitors for 30 min prior to activation: Gö6983 (Sigma-Aldrich), Ly294002 (EMD Chemicals), Rapamycin (EMD Chemicals), 25-hydroxycholesterol (Sigma-Aldrich).

Virus transduction

Truncated forms of SREBP1a or SREBP2 (Supplementary Fig. 1c) were cloned into the retrovirus vector MIGR1 (a gift from Dr. Yukiko Tone and Dr. Masahide Tone, Cedars-Sinai, Los Angeles, USA). Virions were produced using the 293T cell line by transfection using Lipofectamine 2000 (Invitrogen). Purified naive T cells were stimulated in 6-well plates pre-coated with 5 µg/ml of anti-CD3 supplemented with 2 µg/ml of anti-CD28 at 2×10^6 cells/well. At 24 hours, cells were transduced by centrifugation in viral medium plus 8 µg/ml of polybrene and 100 IU/ml of IL-2 at 1200 rpm for 90 min at 37 °C. Medium was then replaced with complete IMDM supplemented with 100 IU/ml of IL-2 and cells were incubated for additional 24 hours. Control cells were subjected to the same stimulation and centrifugation as transduced cells but without virus. After a total of 48 hours of stimulation, cells were harvested and transduction efficiencies were evaluated by GFP expression level on flow cytometry.

RNA interference

Purified T cells were transfected with non-target siRNA, SREBP1 siRNA, or SREBP2 siRNA (Dharmacon) using Nucleofector according to manufacture's protocol (Lonza). Transfected cells were rested for 18 hours at 37 °C and then stimulated with PMA and ionomycin for 5 hours.

Quantitative real-time PCR analysis

RNA was isolated using Trizol (Invitrogen). cDNA was synthesized using iScript cDNA Synthesis Kit (BIO-RAD). qRT-PCR was performed using SYBR Green I Master (Roche) on LightCycler 480 (Roche). Primer sequences are available upon request.

Flow cytometry analysis

7-ADD, anti-mouse CD4 (RM4-5), CD8 (53-6.7), CD25 (PC61.5), CD44 (IM7), Thy1.2 (53-2.1), IL-7R (CD127), KLRG1 (2F1), INF γ (XMG1.2), TNF α (MP6-XT22), and IL-2 (JES6-5H4) were purchased from eBioscience, BioLegend, or BD Biosciences. Active caspase3 and annexin V staining kits were purchased from BD Bioscience. CFDA SE, ER-Tracker Blue-White DPX, and DAPI were purchased from molecular probe. Propidium Iodide was purchased from Calbiochem. Cells were analyzed on LSRII or FACSVerse with FlowJo software (Treestar).

Immunoblot analysis

Whole cell extracts were produced from equivalent number of cells with RIPA buffer (50mM Tris-HCl [pH 8.0], 150mM NaCl, 1% NP40, 0.5% Deoxycholate, 0.1% SDS) supplemented with Calyculin A (Cell Signaling) and protease inhibitor cocktail (Sigma-Aldrich). Samples were separated on a 4-12% Bis-Tris gel and transferred to nitrocellulose. Antibodies for phospho-Akt (S473), Akt1 (5C10), Actin (I-19) were purchased from Santa Cruz Biotechnology. Antibody for HIF1 α was from Cayman Chemical. Antibodies for S6 ribosomal protein (54D2), phospho-S6 (S235/236)(D57.2.2E), Rb (D20), phospho-Rb (Ser807/811), p27Kip1, Cyclin D2 (D52F9), Cyclin D3 (DCS22), MYC, phospho-Zap70 (Tyr319)/Syk (Tyr352)(65E4), phospho-Lck (Tyr505), LAT, phospho-LAT (Tyr191),

p44/42 MAPK (137F5), phospho-p44/42 MAPK (Thr202/Tyr204)(D13.14.4E), AMPK (F6), phospho-AMPK (Thr172)(40H9), phospho-ACC (Ser79) were from Cell Signaling.

Chromatin immunoprecipitation analysis

The method is described elsewhere³⁴. Briefly, 2×10^7 cells were fixed and sonicated. Precleared lysates were incubated overnight at 4 °C with polyclonal anti-SREBP1 and anti-SREBP2, or control rabbit IgG (Cell Signaling). Immunocomplex was collected and real-time PCR was performed as described above. Primer sequences are available upon request.

Gene expression analysis

RNA isolated from littermate control and *Scap*-deficient CD8⁺ T cells and hybridized to Affymetrix GeneChip Mouse Genome 430 2.0 Array. Genes were filtered for differential expression across the samples by requiring an ANOVA p-value less than 0.001. For heatmap display, genes were mean centered, normalized and clustered using a Pearson correlation coefficient metric and pairwise complete-linkage. Pathway enrichment analysis was done with DAVID^{20, 21}. Transcription factor analysis was generated through the use of IPA (Ingenuity Systems, www.ingenuity.com).

Metabolic assays

Cellular ATP levels were measured from the same number of live cells using ATP determination kit according to the manufacture's protocol (Molecular Probes). Background was measured using buffer without cells and subtracted from the values of each cell sample. For mitochondrial mass, cells were incubated with 50 nM MitoTracker green (Molecular Probes) at 37 °C for 30 min. The oxygen consumption rate and extracellular acidification rate were measured with a XF24 analyzer according to the manufacture's recommendations³⁵. Mixing, waiting and measure times were 2, 2, 4 minutes, respectively. Test compounds were obtained from Sigma and injected during the assay at the following final concentration: 0.2 μM oligomycin, 1 μM FCCP, 0.75 μM Rotenone/Myxothiazol. Glucose, glutamine, and lactate concentrations in culture media were measured using bioprofile analyzer (Nova Biomedical).

Cholesterol loading

Cholesterol:MβCD complex containing 40 mg of cholesterol/g was purchased from Sigma (C4951). Treatment concentration was based on cholesterol weight. Cells were activated with or without cholesterol:MβCD in 10% FBS for indicated time.

GC-MS

Purified T cells were collected at *ex vivo* or after 24 hours of activation with plate-bound anti-CD3 in combination with soluble CD28 and washed twice with PBS. Fatty acids extractions and analyses were performed as previously described³⁶ with the modifications described below. Triheptadecanoin (Nu-chek Prep, T-155) was used as the internal standard for fatty acids. Ergosterol (Sigma 17130-U) was used as the internal standard for cholesterol. Internal standards were added to trizol. Cholesterol was extracted from organic phase with petroleum ether after saponification but before acidification and fatty acid extraction.

Extracted cholesterol was trimethylsilylated with BSTFA and TMCS (Sigma 33155-U). Extracted fatty acids were derivatized into their methyl ester form with methanolic boron trifluoride. Derivatized cholesterol and ergosterol were monitored at m/z 456-467 and 468 respectively. Derivatized myristate, palmitate, heptadecanoate, and stearate were monitored at m/z 241-250, 269-278, 284, and 297-306 respectively. Data was collected on an Agilent 5975C MSD connected to an Agilent 7890 Gas Chromatograph with the Phenomenex ZB-MR-1 column. Settings and oven programs are available upon request. Area under the curve quantitation for selected ions was performed on Chemstation software.

LCMV infection, MHC I tetramer and intracellular cytokine staining

Mice were infected intraperitoneally (i.p.) with 2×10^4 plaque forming units (PFU) of LCMV-Armstrong. Virus stocks were prepared and viral titers were quantified as described previously³⁷. Splenocytes were stained directly ex vivo with LCMV-D^bGP₃₃₋₄₁ or LCMV-D^bGP₃₉₆₋₄₀₄ specific tetramers and for surface expression of CD8. MHC tetramers were obtained from the NIH Tetramer Core Facility. To analyze cytokine expression, splenocytes were stimulated for 5 hours with 2 µg/ml of the MHC class I restricted LCMV-GP₃₃₋₄₁ or NP₃₉₆₋₄₀₄ peptide in the presence of 50 U/ml recombinant murine IL-2 (R&D Systems) and 1mg/ml brefeldin A (Sigma-Aldrich). Cells were stained for surface expression of CD8, then fixed, permeabilized and stained with antibodies to TNF, IFN-γ and IL-2 (BioLegend). Flow cytometric analysis was performed using an LSR II or the FACSVerse (Becton Dickinson) and analyzed using FlowJo software (Treestar).

Adoptive cell transfer

CD8⁺ T cells or total T cells were purified from *Scap*^{fl/fl} and control littermate mice or Thy1.1 mice spleens as above. Purified cells were labeled with CFSE and $1-5 \times 10^6$ cells/mouse were injected via retro-orbital into RAG-deficient host or irradiated host 1 day earlier with 600 rad. After 6 days, host spleen and LN cells were analyzed by flow cytometry. For gene expression analysis, Thy1.1⁺ T cells were sorted and real-time PCR was performed as described above.

OVA peptide-IFA immunization

Forty-eight hours after the OT-I cell transfer, mice were either left unimmunized or immunized with total of 300 µg of OVA peptide mixed in IFA into 3 sites in upper back and 3 sites in lower back (n=3 per group). Spleen and lymph nodes were harvested 24 hours later and CD3⁺CD8⁺ cells were analyzed on flow cytometry.

BrdU labeling

BrdU flow kit was purchased from BD Bioscience. BrdU (2 mg/mouse) was injected to *Cd4-Cre/Scap*^{fllox/fllox} and littermate control mice intraperitoneally. Peripheral blood was collected twice a week and mononuclear cells were stained and analyzed on flow cytometry according to manufacture's protocol.

Statistical analysis

Statistical significance was determined with the two-tailed unpaired Student's *t*-test unless otherwise indicated.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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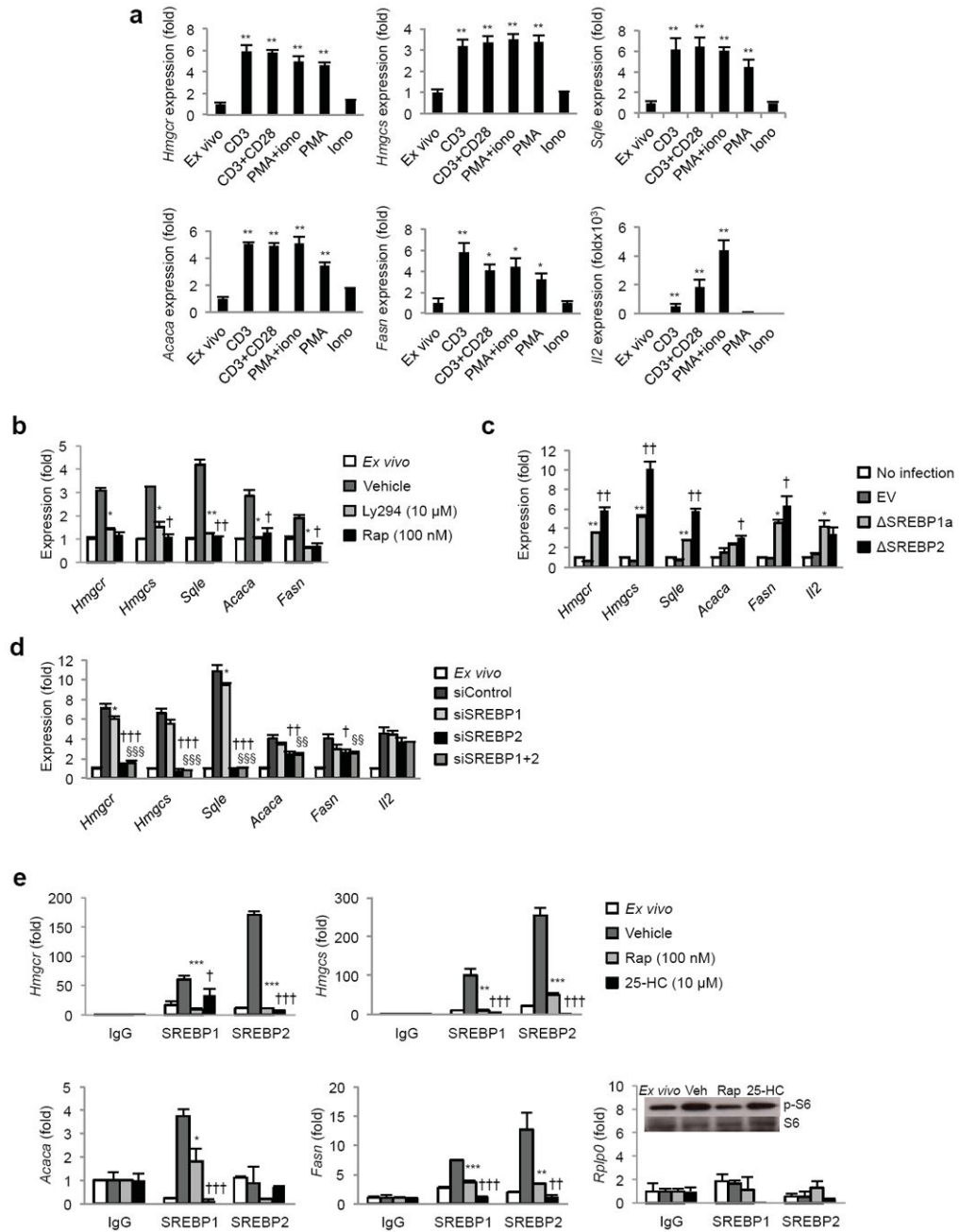


Figure 1. The lipid biosynthetic program of activated T cells is SREBP-dependent and sensitive to the PI3K-mTOR pathway

(a) Real-time PCR analysis of lipogenic genes from murine spleen and LN T cells activated for 6 h with plate-bound CD3+/-CD28, PMA and ionomycin as indicated. (b) Real-time PCR analysis of quiescent or 6 h activated T cells pretreated for 30 min with PI3K inhibitor LY294002 (10 μM) or mTOR inhibitor rapamycin (100 nM) before activation with CD3-28. (c) Real-time PCR analysis of lipogenic genes in activated T cells transduced with active SREBP (ΔSREBP1a or ΔSREBP2) or empty vector (EV) for 24 h. * denotes comparisons between empty vector and ΔSREBP1a, † denotes comparisons between empty vector and

Δ SREBP2. Number of symbols indicates the p value. **(d)** Real-time PCR analysis of quiescent or T cells transfected with siSREBP1 alone, siSREBP2 alone, siSREBP1 and siSREBP2 or non-target control (siControl) and then activated with PMA-iono for 5 h. * denotes comparisons between siControl and siSREBP1, † denotes comparisons between siControl and siSREBP2, § denotes comparisons between siControl and siSREBP1+2. **(e)** Chromatin Immunoprecipitation analysis of SREBP1 and SREBP2 at the promoters of indicated genes in splenocytes *ex vivo* or 4 hours after activation with PMA. In addition, some cultures were pretreated for 30 minutes with rapamycin (100 nM) or 25-hydroxycholesterol (25-HC, 10 μ M) before activation. Data is normalized to input and expressed relative to IgG control. *Rplp0*: non-SREBP target gene control. * Denotes comparisons between vehicle and rapamycin treated samples, † denotes comparisons between vehicle and 25-HC treated samples. Inset: Immunoblot of phospho and total S6 from whole cell lysates to confirm rapamycin function. For all experiments, data are presented as a mean of triplicates with standard deviation, and representative of at least three experiments. *p<0.05, **p<0.01, ***p<0.001 (statistical difference between other experimental groups is also noted by the number of other symbols, two-tailed unpaired Student's *t*-test).

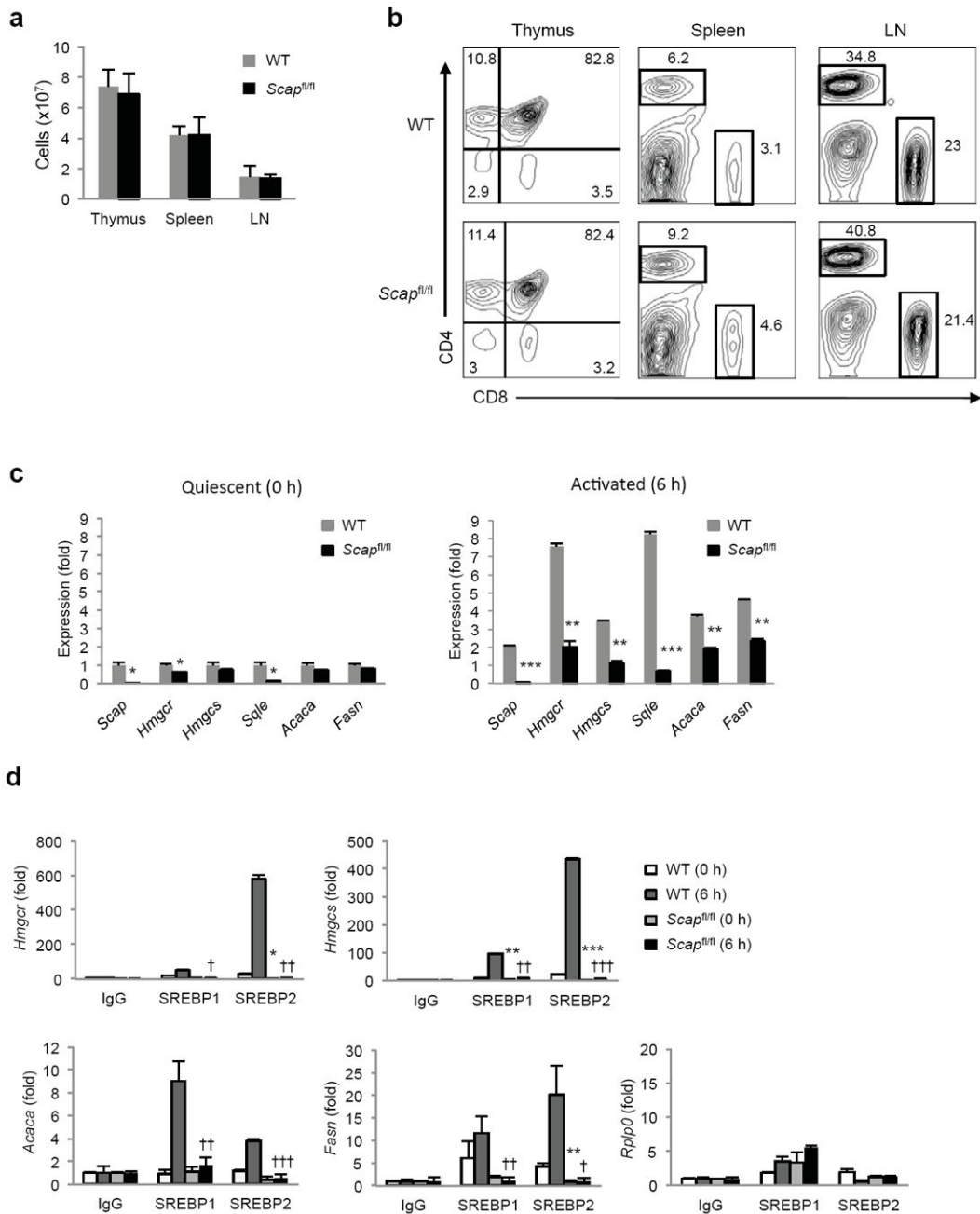


Figure 2. Deletion of *Scap* inhibits SREBP activity but does not affect T cell homeostasis (a) Total cell number in thymus, spleen and lymph nodes (LN) from *Cd4-Cre-Scap^{fl/fl}* mice (designated *Scap^{fl/fl}* herein) and littermate controls (WT). Data is expressed as mean with standard deviation. (b) Flow cytometric analysis of CD4⁺ and CD8⁺ T cells in thymus, spleen and LN from *Scap^{fl/fl}* and littermate control mice. Relative frequency indicated in plots. (c) Real-time PCR analysis of control and *Scap^{fl/fl}* CD8⁺ T cells ex vivo or activated for 6 h with anti-CD3-28. (d) Chromatin immunoprecipitation analysis of SREBP1 and SREBP2 at the promoters of indicated genes in T cells ex vivo or 6 hours after activation (anti-CD3-28). Data are normalized to input and plotted relative to IgG control. *Rplp0*: non-

SREBP target gene control. * denotes comparisons between WT T=0 and *Scap*^{fl/fl} T=0, † denotes comparisons between WT T=6 and *Scap*^{fl/fl} T=6. Data are mean of triplicates with standard deviation (**c,d**). *p<0.05, **p<0.01, ***p<0.001 (statistical difference between other experimental groups is also noted by the number of other symbols, two-tailed unpaired Student's *t*-test). Data are representative of four independent experiments with three mice per group (**a,b**), or three independent experiments (**c,d**).

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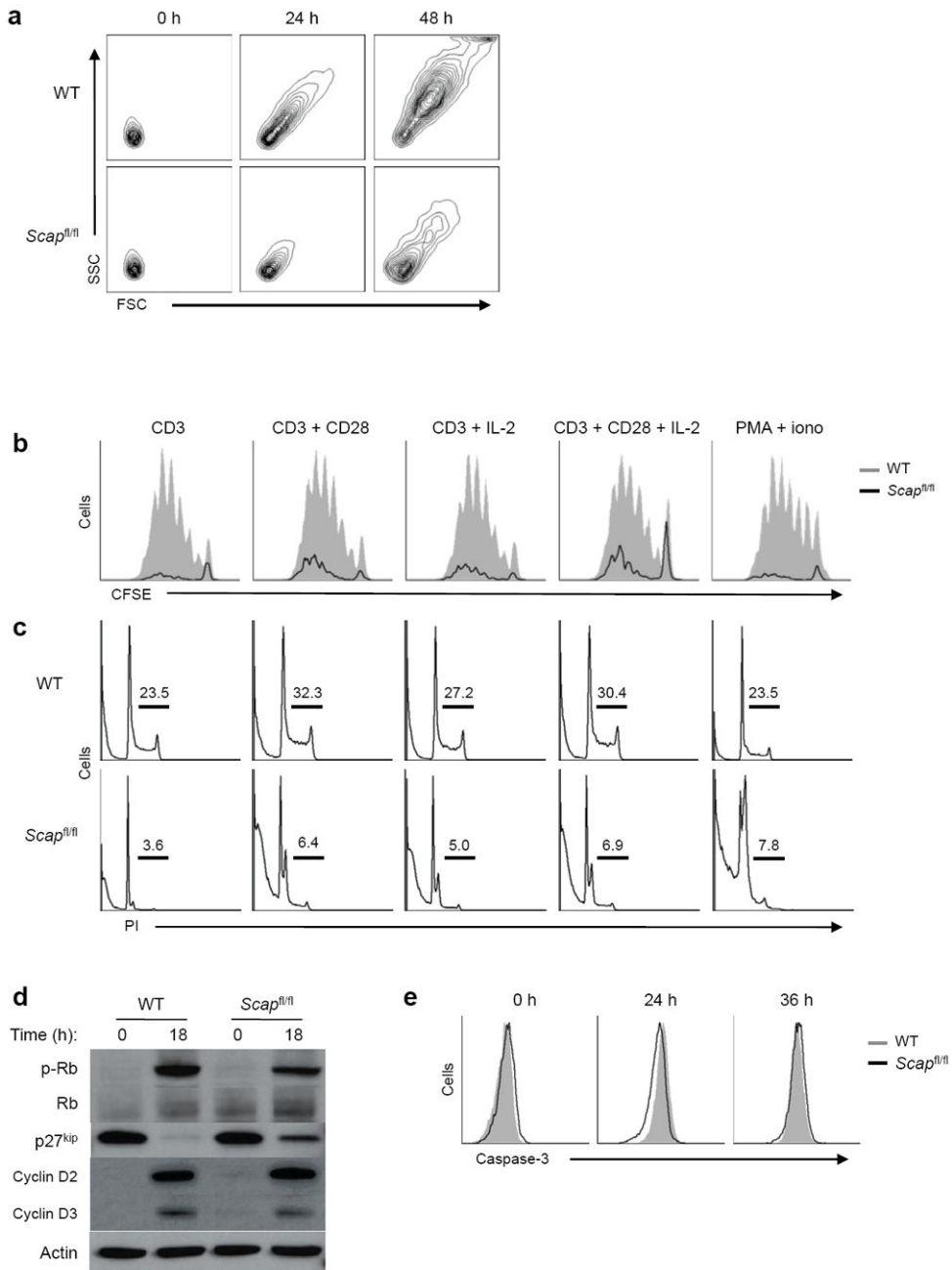


Figure 3. SREBP activity influences CD8⁺ T cell growth and proliferation

(a) Forward and side scatter plots of DAPI-negative (live) WT and *Scap*^{fl/fl} CD8⁺ T cells activated with anti-CD3-28 for indicated time. (b) CFSE dilution of DAPI negative (live) WT and *Scap*^{fl/fl} CD8⁺ T cells activated for 72 hours with indicated mitogens. (c) Flow cytometric analysis of DNA content from WT and *Scap*^{fl/fl} CD8⁺ T cells activated for 48 hours with indicated stimuli. Cells were stained for DNA content with propidium iodide. Frequency of S and G2/M phase indicated in plots. (d) Immunoblots of G1 associated cell cycle proteins from whole cell lysates of quiescent or 18 h activated WT and *Scap*^{fl/fl} CD8⁺ T cells with anti-CD3-28. (e) Flow cytometric analysis of intracellular cleaved caspase-3 in

WT and *Scap^{fl/fl}* CD8⁺ T cells activated with anti-CD3-28 for indicated time. Data are representative of three (**a**), or two (**b,c,d,e**) independent experiments.

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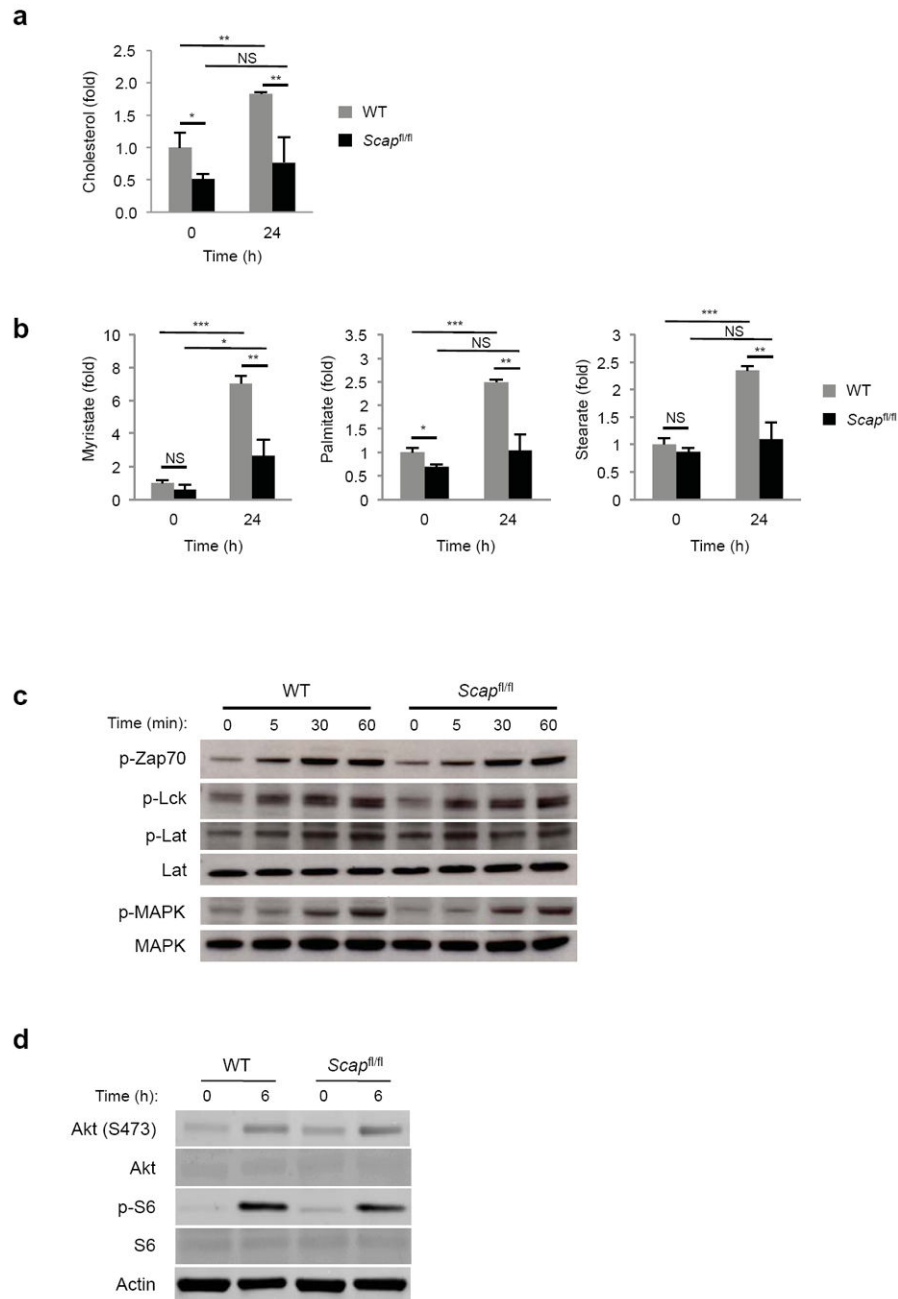


Figure 4. Loss of SREBP signaling impacts lipid homeostasis but does not perturb proximal TCR signaling

(a,b) GC-MS determination of cholesterol (a) or indicated long-chain fatty acid (b) content of control and *Scap*^{fl/fl} T cells *ex vivo* or 24 h after activation with anti-CD3-28. Data are normalized to cell numbers and plotted relative to control *ex vivo* (WT 0 h). Data is expressed as mean of triplicates with standard deviations. (c) Immunoblot analysis of WT and *Scap*^{fl/fl} lymphocytes *ex vivo* or after stimulation with 1 μ g/ml of soluble anti-CD3e for the indicated time. Whole cell lysates were blotted for indicated total and phosphorylated proteins. (d) Immunoblot analysis of the Akt pathway from WT and *Scap*^{fl/fl} CD8⁺ T cells

ex vivo or after 6 h of activation with anti-CD3-28. NS: not significant, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ (two-tailed unpaired Student's *t*-test).

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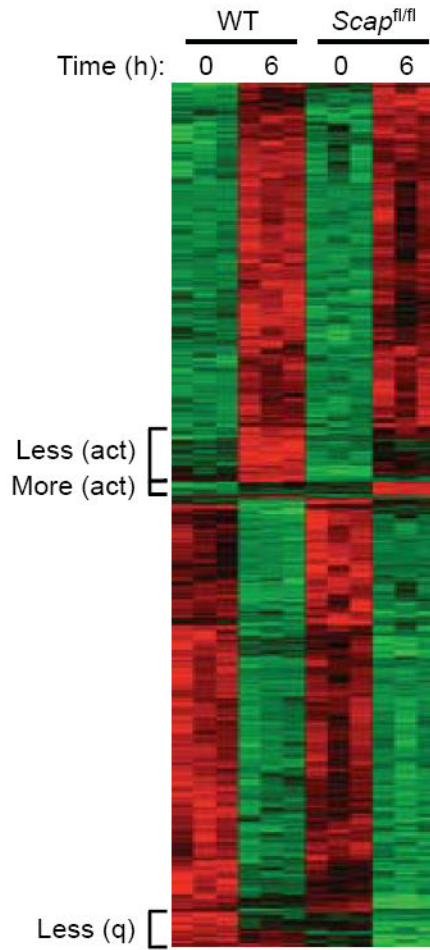


Figure 5. SREBP activity influences a transcriptional program related to lipid and RNA metabolism

Heat map of genes expressed in purified quiescent or 6 h stimulated (anti-CD3-28) WT and *Scap*^{fl/fl} CD8⁺ T cells (all changes in gene expression met criteria of $p < 0.001$ to be considered as significant). Less (act): less induced in activated *Scap*^{fl/fl} CD8⁺ T cells. More (act): more induced in activated *Scap*^{fl/fl} CD8⁺ T cells. Less (q): less induced in quiescent *Scap*^{fl/fl} CD8⁺ T cells.

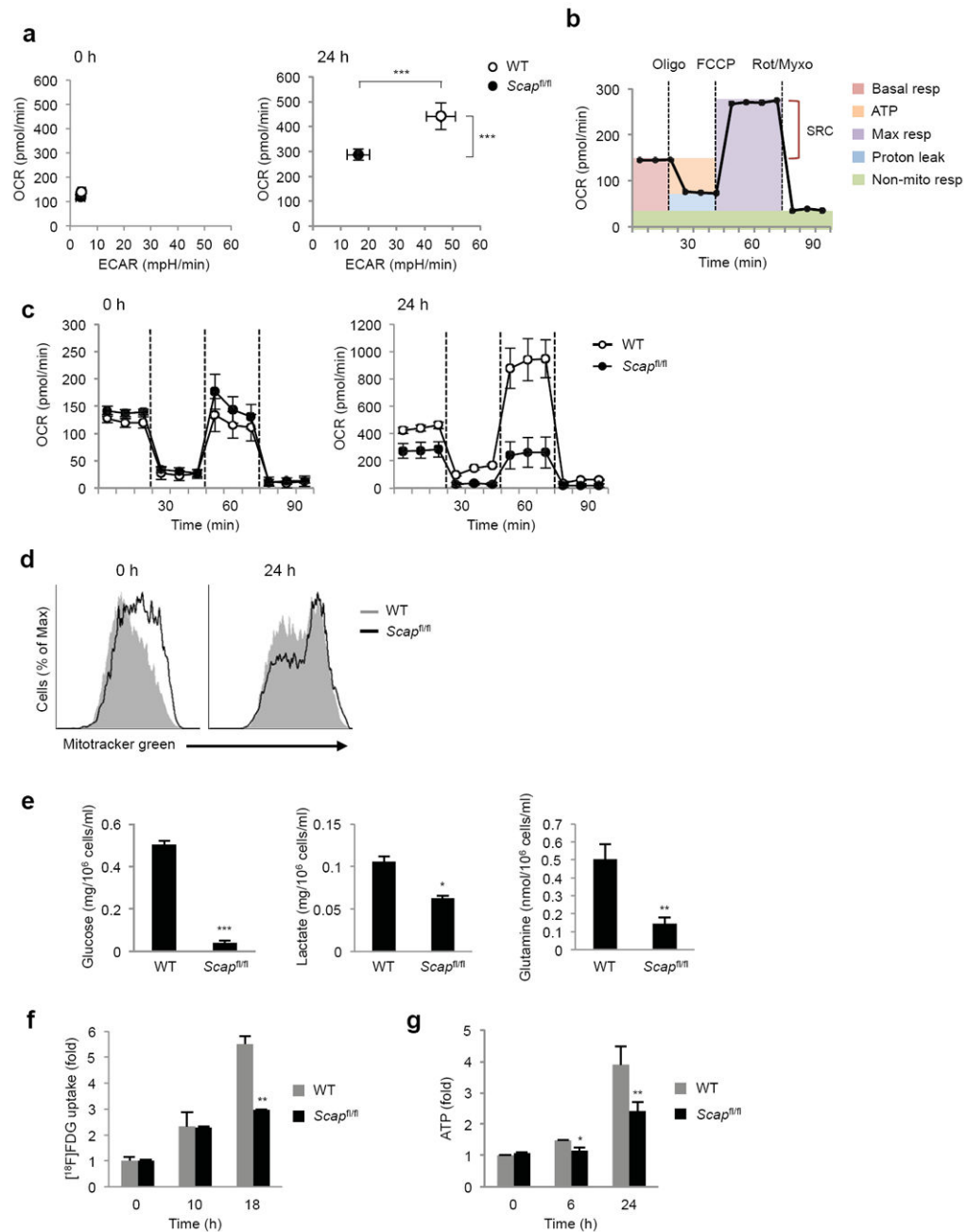


Figure 6. SREBP signaling is required for metabolic reprogramming of activated CD8⁺ T cells
(a) Basal oxygen consumption rate (OCR) and extracellular acidification rate (ECAR) of quiescent and 24 h stimulated WT and *Scap^{fl/fl}* CD8⁺ T cells. **(b)** A schematic of OCR in basal condition and in response to sequential treatment with oligomycin (oligo: ATPase inhibitor), FCCP (uncoupling agent) and rotenone-myxothiazol (Rot-Myxo: electron transport chain inhibitor). Basal resp: basal respiration, ATP: ATP production, Max resp: maximal respiration, Non-mito resp: non-mitochondrial respiration, SRC: spare respiratory capacity **(c)** OCR of 24 h activated WT and *Scap^{fl/fl}* CD8⁺ T cells in basal state and in response to sequential treatment with oligomycin, FCCP and rotenone-myxothiazol. **(d)**

Mitotracker staining of WT and *Scap*^{fl/fl} CD8⁺ T cells *ex vivo* and after 24h of activation with anti-CD3-28. (e) Glucose and glutamine consumption, and lactate production measurement in spent media from activated WT and *Scap*^{fl/fl} CD8⁺ T cell activated for 24 h with anti-CD3-28. (f) 2-deoxy-2-(¹⁸F)fluoro-D-glucose (FDG) uptake assay. CD8⁺ T cells were activated with anti-CD3-28 for indicated time. Cultures were pulsed with [¹⁸F]FDG for 60 min before analysis. (g) Cellular ATP levels in quiescent, 6 or 24 h activated WT and *Scap*^{fl/fl} CD8⁺ T cells (anti-CD3-28). Data are mean of triplicates with standard deviation and representative of two (c) or three (d) independent experiments. *p<0.05, **p<0.01, ***p<0.001 (two-tailed unpaired Student's *t*-test).

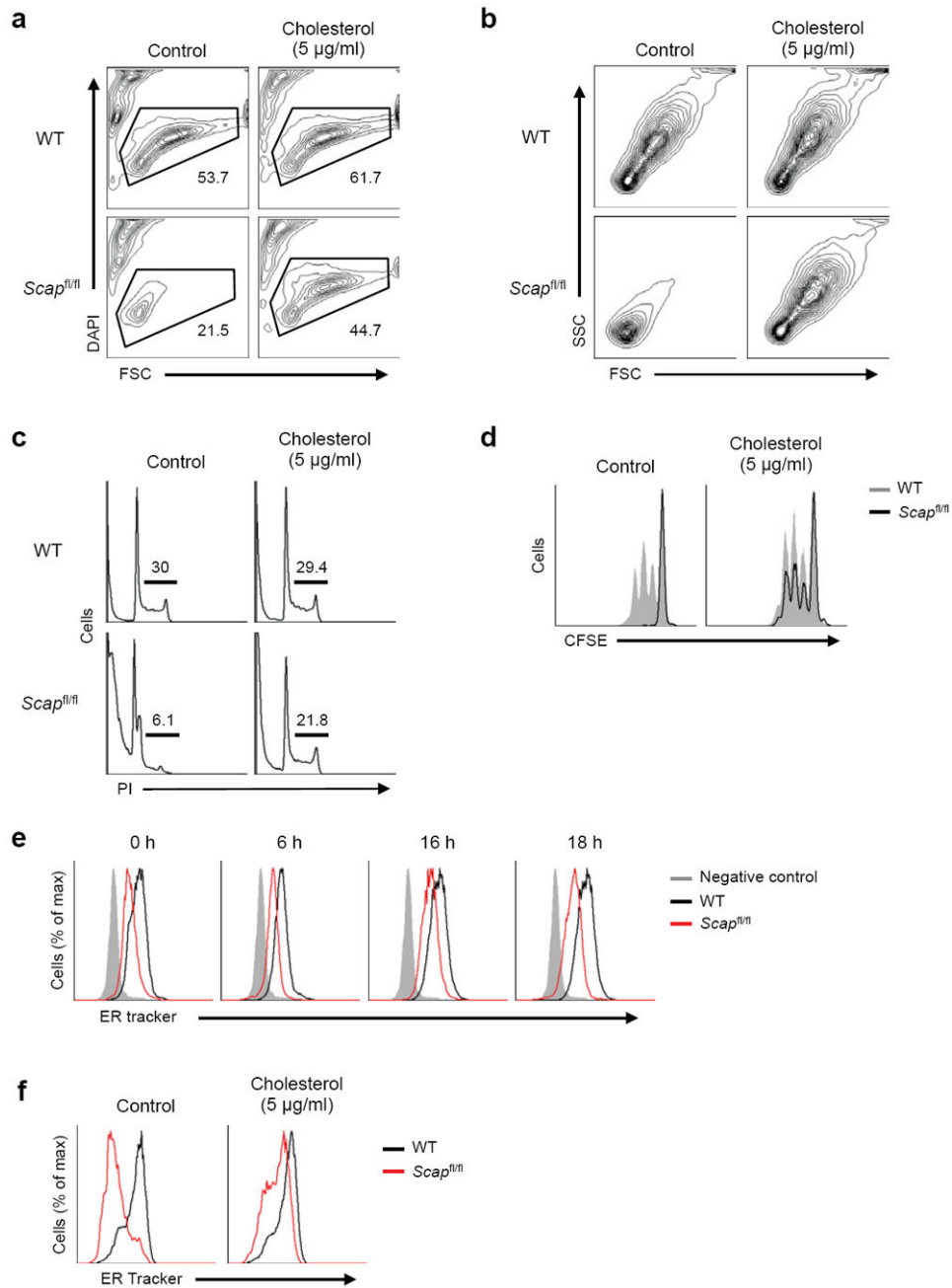


Figure 7. Addition of cholesterol rescues growth and proliferation impairment of *Scap*-deficient CD8⁺ T cells

(a) Viability of WT and *Scap^{fl/fl}* CD8⁺ T cell activated with anti-CD3-28 for 48 h with or without 5 µg/ml of MβCD conjugated cholesterol in 10% FBS supplemented media. DAPI-negative (live) cell populations are gated and the frequencies are indicated in plots. (b) Forward and side scatter plots of DAPI-negative (live) WT and *Scap^{fl/fl}* CD8⁺ T cell activated with anti-CD3-28 for 48 h with or without 5 µg/ml of MβCD-cholesterol in 10% FBS supplemented media. (c) Flow cytometric analysis of DNA content from WT and *Scap^{fl/fl}* CD8⁺ T cells activated for 48 hours with or without 5 µg/ml of MβCD-cholesterol

in 10% FBS supplemented media. Cells were stained for DNA content with propidium iodide. Frequency of S and G2/M phase indicated in plots. **(d)** CFSE dilution of DAPI negative (live) WT and *Scap^{fl/fl}* CD8⁺ T cells activated for 48 hours with or without 5 µg/ml of MβCD-cholesterol in 10% FBS supplemented media. **(e)** ER-tracker staining of WT and *Scap^{fl/fl}* CD8⁺ T cells ex vivo or after activation with anti-CD3-28 for indicated time. **(f)** ER-tracker staining of WT and *Scap^{fl/fl}* CD8⁺ T cells activated for 48 hours with or without 5 µg/ml of MβCD-cholesterol in 10% FBS supplemented media.

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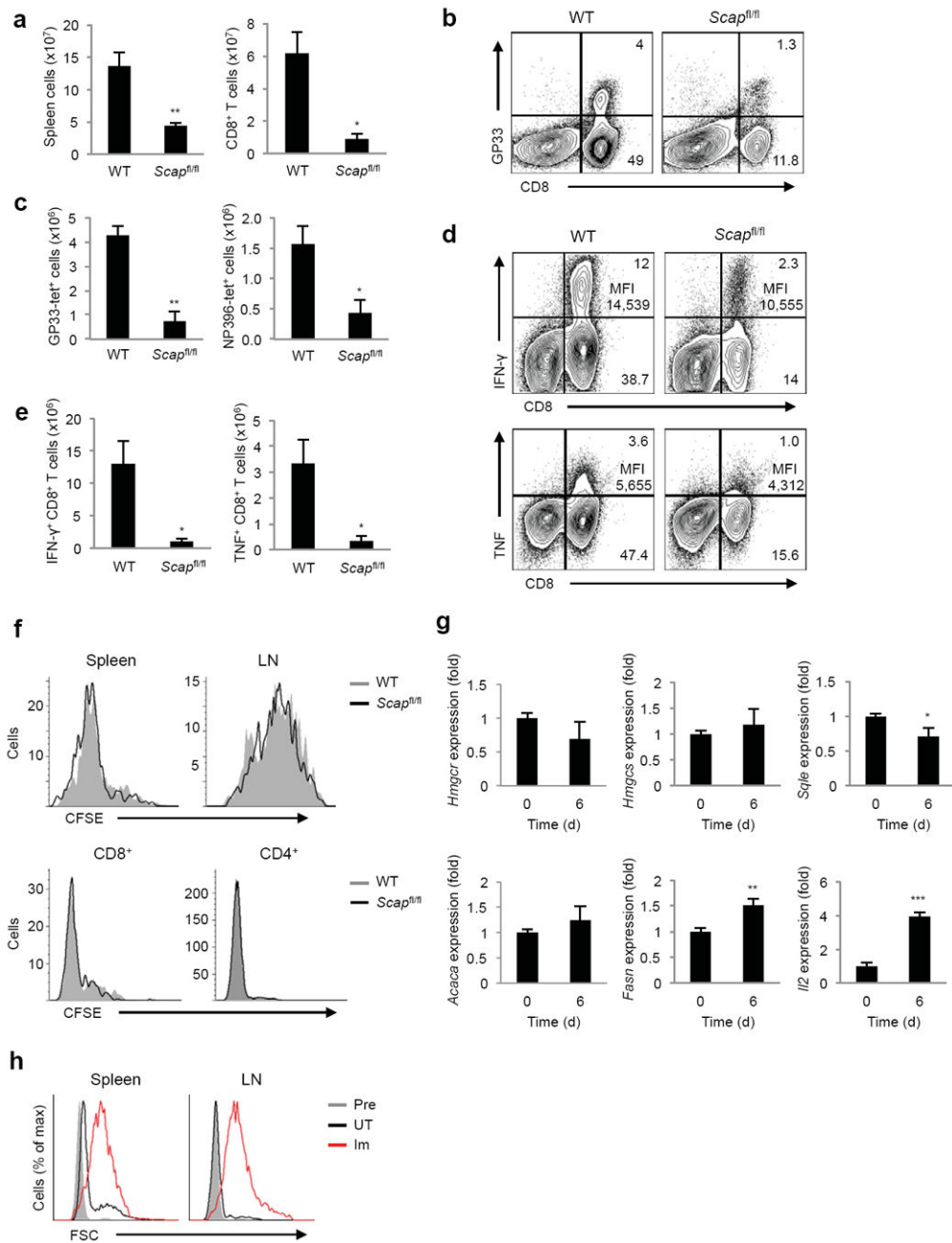


Figure 8. Loss of SREBP activity impairs clonal expansion of antigen specific effector CD8⁺ T cells, but does not perturb homeostatic proliferation

(a) Absolute numbers of splenocytes and CD8⁺ T cells from spleen harvested on day 8 of LCMV Armstrong (Arm) infection. (b) Frequency of D^bGP33-tetramer specific CD8⁺ T cells from spleen harvested on day 8 of LCMV-Arm infection. Frequency of cells marked in quadrants of FACS plots. (c) Absolute numbers of CD8⁺D^bGP33-tetramer⁺ cells and CD8⁺NP396-tetramer⁺ cells from spleen harvested on day 8 of LCMV-Arm infection. (d,e) Frequency (d) and absolute number (e) of IFN- γ or TNF producing WT and *Scap^{fl/fl}* CD8⁺ T cells harvested on day 8 of LCMV-Arm infection and stimulated with GP33-peptide for

5h *in vitro*. Frequency and mean fluorescent intensity (MFI) of cytokine producing CD8⁺ T cells indicated in right upper quadrant of FACS plots. (f) CFSE dilution gated on Thy1.2⁺ CD8⁺(top), and CD3⁺CD4⁺ or CD3⁺CD8⁺ (bottom) cells on day 6 after adoptive transfer into RAG-deficient Thy1.1⁺ mice (top) or irradiated littermate WT mice (bottom). (g) Real-time PCR analysis of Thy1.1⁺ T cells adoptively transferred into RAG-deficient Thy1.2⁺ mice and sorted from spleens and LNs after 6 days. (h) Forward scatter plots of OT-I cells. Cells were adoptively transferred into RAG-deficient mice and either left untreated (UT) or immunized (Im) with OVA/IFA subcutaneously. Twenty-four hours later, cells from spleens and LNs were stained as described in Methods and analyzed on flow cytometry. Pre: pre-transfer. *p<0.05, **p<0.01 (two-tailed unpaired Student's *t*-test).

Table 1
Pathway enrichment analysis of genes less induced in 6 h activated *Scap^{fl/fl}* versus WT CD8⁺ T cells using DAVID (NIAID)

The number of genes enriched in each pathway (designated “count”) and associated p values are indicated.

Enrichment cluster 1: Score 7.09	Count	P Value
sterol biosynthetic process	10	6.20×10^{-14}
sterol metabolic process	12	6.10×10^{-13}
steroid biosynthetic process	11	8.50×10^{-12}
cholesterol biosynthetic process	8	3.90×10^{-11}
steroid metabolic process	13	1.30×10^{-10}
cholesterol metabolic process	10	2.30×10^{-10}
lipid biosynthetic process	14	8.90×10^{-9}
isoprenoid biosynthetic process	4	2.70×10^{-4}
isoprenoid metabolic process	4	2.90×10^{-3}
oxidation reduction	11	5.40×10^{-3}
Enrichment cluster 2: Score 4.33	Count	P Value
nuclear lumen	20	2.90×10^{-7}
intracellular organelle lumen	22	7.00×10^{-7}
organelle lumen	22	7.30×10^{-7}
nucleolus	12	1.20×10^{-6}
membrane-enclosed lumen	22	1.20×10^{-6}
nucleoplasm	10	6.60×10^{-3}
intracellular non-membrane-bounded organelle	16	1.20×10^{-1}
non-membrane-bounded organelle	16	1.20×10^{-1}
Enrichment cluster 3: Score 2.75	Count	P Value
ribonucleoprotein complex biogenesis	8	1.40×10^{-5}
ribosome biogenesis	7	4.70×10^{-5}
rRNA processing	5	8.90×10^{-4}
rRNA metabolic process	5	9.40×10^{-4}
RNA processing	9	3.80×10^{-3}
ribonucleoprotein complex	9	4.50×10^{-3}
ncRNA processing	5	1.30×10^{-2}
ncRNA metabolic process	5	3.00×10^{-2}
Enrichment cluster 4: Score 2.72	Count	P Value
RNA binding	16	6.90×10^{-6}
RNA recognition motif, RNP-1	7	7.10×10^{-4}
Nucleotide-binding, alpha-beta plait	7	1.10×10^{-3}

Enrichment cluster 1: Score 7.09	Count	P Value
mRNA metabolic process	6	3.10×10^{-2}
mRNA splicing	4	8.40×10^{-2}
RNA splicing	4	1.10×10^{-1}
mRNA processing	4	1.40×10^{-1}
mRNA processing	4	2.00×10^{-1}

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Table 2

Predicted transcription factors for the genes less induced in 6 h activated *Scap*^{fl/fl} versus WT CD8⁺ T cells.

Transcription Factor	P value	FDR
CREBBP	2.47×10^{-9}	4.55×10^{-3}
SREBF2	2.48×10^{-9}	6.82×10^{-3}
SREBF1	2.65×10^{-8}	9.09×10^{-3}
NFYA	3.86×10^{-5}	1.14×10^{-2}
YY1	6.69×10^{-4}	1.36×10^{-2}

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