Title: Single-cell lineage trajectories and chromatin regulators that initialize antiviral CD8 T cell ontogeny

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Summary

Individual naive CD8 T cells activated in lymphoid organs differentiate into functionally diverse and anatomically distributed T cell phylogenies in response to intracellular microbes. During infections that resolve rapidily, including live viral vaccines[1], distinct effector (T_{EFF}) and memory (T_{MEM}) cell populations develop that ensure long term immunity[2]. During chronic infections, responding cells progressively become dysfunctional and "exhaust"[3]. A diverse taxonomy of T_{EFF}, T_{MEM} and exhausted (T_{EX}) CD8 T cell populations is known, but the initial developmental basis of this phenotypic variation remains unclear[4-10]. Here, we defined single-cell trajectories and identified chromatin regulators that establish antiviral CD8 T cell heterogeneity using unsupervised analyses of single-cell RNA dynamics[11-13] and an *in vivo* RNAi screen[14]. Activated naive cells differentiate linearly into uncommitted effector-memory progenitor (EMP) cells, which initially branch into an analogous manifold during either acute or chronic infection. Disparate RNA velocities in single EMP cells initiate divergence into stem, circulating, and tissue-resident memory lineages that generate diverse T_{MEM} and T_{EX} precursor states in specific developmental orders. Interleukin-2 receptor (IL-2R) signals are essential for formation and transcriptional heterogeneity of EMP cells, and promote trajectories toward T_{EFF} rather than T_{EX} states. Nucleosome remodelers *Smarca4* and *Chd7* differentially promote transcription that delineates divergent T_{MEM} lineages before cooperatively driving terminal T_{EFF} cell differentiation. Thus, the lineage architecture is established by specific chromatin regulators that stabilize diverging transcription in uncommitted progenitors.

Main text

To clarify the initial origins of T cell memory we generated longitudinal single-cell RNA-sequencing (scRNA-seq) datasets and used unsupervised methods to map single-cell trajectories that developed from naive CD8 T cells specific for *Lymphocytic choriomeningitis virus* (LCMV) early after infection of wildtype mice with strains that cause either an acute (Armstrong, $LCMV_{Arm}$), or chronic (Clone 13, $LCMV_{Cl13}$) infection (fig S1A-B) [15]. On days 5 and 8 post infection (pi), clonal TCR transgenic P14 cells that had been adoptively transferred and endogenous polyclonal MHC-I tetramter-reactive CD8 T cells ($GP33^+$), which both recognize the LCMV epitope GP_{33-41} in MHC $H\text{-}2D^b$, were isolated from the spleens of the same host mice, and libraries were generated in parallel with fresh naive CD8 T cells purified from separate P14 mice. Individual naive cells are recruited into the response over the first ~ 3 days following primary infection [16]. Due to this asynchrony, we anticipated the time-series sampling would encompass multiple developmental states that compose initial antiviral CD8 T cell ontogeny.

Dimensionality reduction and Louvain cluster extraction of cells was performed on all samples simultaneously using similar numbers of randomly sampled cells from each experimental group to limit potential biological biases arising from changes in subset compositions at different time points, and the data are represented in the two-dimensional PAGA initialized force-directed (FA) embedding (**Fig 1A** and fig S1B) [17,18]. Partition-based graph abstraction (PAGA) inferred single-cell paths based on correlations between clusters [13], which were numbered according to pseudotime (P0-P10) to define a potential developmental order (**Fig 1A-B,** Table S1). As expected, naive cells (P0) clustered apart from all activated cells, which separated into multiple clusters (P1-P10) (**Fig 1A** and fig S1C-D) and the pseudotime arrangement correctly predicted the actual time dependent emergence of cells in specific clusters (naive, vs days 5 and 8 pi) (**Fig 1C,** S1C-D and Table S1). The distribution of P14 and $GP33^+$ CD8 T cells among the clusters was similar in $LCMV_{Arm}$-infected mice (fig S1D and E). P14 cells isolated from $LCMV_{Arm}$ and $LCMV_{Cl13}$ infected hosts were distributed in similar clusters on day 5 pi, but contributed differentially to clusters on day 8 pi (**Fig 1C** and fig S2D and E). Cellular identities of the clusters were imputed using gene set enrichment analysis (GSEA) and "subtractive" gene expression signatures extracted from published bulk-RNA-seq data derived from phenotypically defined CD8 T cell subsets [19,20] (Table S2, https://github.com/TCellResearchTeam/T_cell_signature_Reference). All major T_{EFF}, T_{MEM}, T_{EX} and naive cell signatures were strongly enriched (p-val <0.05, NES) in at least one Louvain cluster (**Fig 1D** and fig S1 J-M), and demonstrated that cells corresponding to all mature T_{EFF},

T_{MEM} and T_{EX} cell gene expression states arise within 8 days following acute or chronic LCMV infection. The PAGA-inferred paths between these states facilitated precisely defining developmentally regulated gene expression at the single-cell level, which extends previous longitudinal studies of bulk populations during acute infection [21].

Naive CD8 T cells differentiate along a linear path into common effector and memory progenitor (EMP) cells

The unsupervised approach clarified the initial developmental relationship of T_{MEM}, T_{EFF}, and T_{EX} cells in an ubiased fashion (**Fig 1B**). Naive cells (P0) were connected to cells in cluster P2, via activated intermediates (P1 cells, **Fig 1E** and Table S1). P2 cells were positively enriched with gene expression of recent TCR stimulation (48h Act up, p = 0.004, NES = 2.1). P4 cells were negatively enriched with this signature (fig S1K), and both GSEA and pseudotime indicated P4 cells were more developmentally advanced than those in P2, and were therefore downstream (fig S1J, Best clusters 2, 8 and 10). Thus, activated naive cells appeared to initially develop along a linear pathway into P2 cells.

Transcriptionally heterogeneous cell clusters on day 5 emerged from P2 cells, which strongly expressed *Il2ra* (encodes CD25/IL-2Rα a subunit of the trimeric interleukin-2 receptor that initiates high-affinity IL-2 binding[22,23]) and were positively enriched with signatures of both central memory (T_{CM}) and naive cells, but not those of mature effector memory (T_{EM}), memory stem (T_{SCM}), resident memory (T_{RM}) or terminal effector (TE) cells (**Fig 1D** and fig S1L). P2 cells highly expressed a mixture of genes encoding TFs whose cognate motifs are enriched within *cis*-acting regions that gain *de novo* chromatin accessibility during primary TCR stimulation (*Runx3*, *Batf*, *Irf4, Prdm1*, *Klf2*), and that are essential for both T_{EFF} and T_{MEM} cell development[24,25](fig S1G and H). In addition, genes encoding multiple regulatory factors whose expression is highly differential in established mature TE/T_{EM} (*Tbx21*, *Zeb2*, *Id2, Prdm1*), T_{CM}/T_{SCM}/T_{EX}^{prog1} (*Tcf7*, *Zeb1*, *Bach2*, *Id3*), T_{RM} (*Hmmr, Aurkb, Prdm1*) and T_{EX}^{term} (*Tox*, *Lag3, Cd160*) populations[26,27] were coordinately expressed at intermediate levels in P2 cells (fig S1G and H). These “lineage-specific” genes were significantly upregulated or downregulated in cells from clusters at the distal tips of the paths (P10, P7, P3 and P8) compared to P2 cells (**Fig 1E**, fig S1G and H and Table S1), implying P2 cells promiscuously express regulatory factors that become progressively lineage-restricted. Flow cytometry confirmed that activated naive cells exhibited uniform behavior while undergoing extensive cell division, upregulating CD25 and maintaining expression of T_{CM} cell attributes (CD27 and Tcf1 expression), before developing

phenotypic features of more mature T_{EFF} cells (e.g., high Blimp1-YFP and KLRG1 expression) (**Fig 1F**). These divergent subsets emerged near the final detectable cell division from cells highly expressing both CD25 and the naive and T_{CM}-associated TF Tcf1 (fig S1I). Thus, naive cells initially differentiate in a linear path into cells that manifest multilineage gene expression, a hallmark of multipotency and cells undergoing lineage-choice [28,29]. On this basis, we classified cells in cluster P2 as common effector/memory progenitor (EMP^{P2}) cells.

Disparate RNA velocities develop in individual EMP cells and initiate a branched manifold that establishes T_{MEM} and T_{EX} cell diversity

Strong connections of EMP^{P2} cells with clusters arranged immediately downstream implied the initial branchpoints of four developmental paths (**Fig 1B**). To define the trajectories of cells from each cluster in the PAGA-inferred architecture, their future states were modeled using RNA velocity[11,12] (**Fig 2A**). Nascent RNA expression precedes accumulation of their mature mRNAs by several hours, and RNA velocity describes the rates at which cells are transitioning into new states based on the gene-wise ratios in expression of nascent (i.e. unspliced) to mature (i.e., spliced) mRNAs genome-wide. Streamline plots after UMAP embedding of all samples from each infection depicted transition probability data derived from the grid average RNA velocities between single-cell clusters (**Fig 2A** and fig S2A), and defined future cell states in the lineage architecture (**Fig 2B** and **2C** fig S2A). Strongly divergent RNA velocities in P2 and P5 cells confirmed they were developmental roots in each infectious context (fig S2B). The RNA velocities of signature genes associated with multiple distinct CD8 T cell states (e.g., naive and T_{SCM} cells: *Id3, Tcf7* and *Sell*; and TE cells: *Prdm1*, *Id2*, *Tbx21* and *Zeb2*) were all positive in EMP^{P2} cells during $LCMV_{Arm}$ infection (**Fig 2E**, Table S5). This indicates that multilineage gene transcription in rapidly dividing EMP^{P2} cells establishes the transition potential into diverse future cell states, prior to developmental branching and mature lineage-specific mRNA expression. Conversely, cells in cluster P10 during $LCMV_{Arm}$ infection, and clusters P7 and P8 during $LCMV_{Cl13}$ infection, lacked RNA velocities into other clusters indicating they were terminal states in the analysis (fig S2C).

Trajectory 1 (T1: P2->P4->P9->P10) only formed in $LCMV_{Arm}$-infected mice and defined formation of cells enriched with signatures of bulk T_{EM} cells (effector memory precursors, pT_{EM}^{P4}), classical $KLRG1^{lo}$ $CD127^{lo}$ early effector cells (EE^{P9}) and $KLRG1^{hi}$ $CD127^{lo}$ terminal effector (TE) cells (TE^{P10}) (**Fig 2C-D**, fig S1K and M). RNA velocity-derived transition probabilities indicated most EE^{P9} cells proceed toward TE^{P10} cells (**Fig 2B**). Positive *Tbx21*

RNA velocity was sustained and *Zeb2* velocity accelerated in EE^{P9} cells, followed by increased expression of mature *Tbx21* and *Zeb2* mRNAs in TE^{P10} cells, whose velocities continued to increase in TE^{P10} cells[30-32](**Figs 2D-E** and fig S2D and Table S5). Thus, the initial developmental order of T_{CIRC} cells during acute LCMV infection is T_{CM} to T_{EM} to TE cells.

Trajectory 2 (T2: P2>P5>P3) is likely a main source of T_{RM} precursor cells (pT_{RM}^{P3}) during $LCMV_{Arm}$ infection, and exhausted progenitor 2 ($T_{EX}^{prog2\text{-}P3}$) cells[33] during $LCMV_{Cl13}$ infection. In $LCMV_{Arm}$-infected hosts, P5 cells were positively enriched with signatures of bulk $KLRG1^{hi}CD127^{hi}$ "double positive" (DP) effector and T_{RM} cells, and were classified as DP^{P5} cells (**Fig 1D** and fig S1K). DP^{P5} cells diverged into pT_{RM}^{P3} cells, pT_{EM}^{P4} cells, and cells enriched with the bulk signature of classical $KLRG1^{lo}CD127^{hi}$ memory precursor (MP) cells (MP^{P6}) (**Figs 1B and 2A-C** and fig S1K-L). Runx3-dependent gene expression was positively enriched in DP^{P5} and MP^{P6} (fig S1K and S2F), consistent with the requirement of Runx3 for development of these cell states[25,34], and divergence into pT_{RM}^{P3} cells correlated with increased expression of the reprsentative T_{RM} signature gene *Hmmr* (**Fig 2B-D**, and fig S1H). In $LCMV_{Cl13}$-infected hosts P5 cells were also enriched with signatures of DP and T_{RM} cells, however, they were classified as $T_{EX}^{prog2\text{-}P5}$ cells because they were positively enriched with the signatures of both T_{EX}^{prog2} and T_{EX} cells (**Fig 1D** and fig S1L-M). In addition, cluster P3 cells were enriched with both T_{RM} and T_{EX}^{prog2} signatures, and were classified as $T_{EX}^{prog2\text{-}P3}$ cells, but were distinct from $T_{EX}^{prog2\text{-}P5}$ cells because they lacked TCR-stimulated gene expression (fig S1K-L). The T_{EX} signature was not strongly enriched in either DP^{P5} or pT_{RM}^{P3} cells from $LCMV_{Arm}$-infected mice, which confirmed their distinction from homologous cells during chronic infection which were and that strongly expressed *Tox*[35-37] (**Figs 1D and 2E**, and fig S1M).

Trajectory 3 (T3: P2>P6>P8) explained initial development of intermediately exhausted (T_{EX}^{int}) and terminally exhausted cells (T_{EX}^{term}) during $LCMV_{Cl13}$ infection, and formation of classical MP cells during $LCMV_{Arm}$ infection (**Fig 2C**). In $LCMV_{Arm}$-infected hosts, MP^{P6} cells were enriched with MP and EE cell signatures and had transition potential into EE^{P9} cells, providing an alternative conduit into the T1 trajectory (**Fig 2A-C,** fig S1K-M), and demonstrated classical MP cells are most likely distinct from precursors of T_{SCM} (see below). P6 cells from $LCMV_{Cl13}$-infected were classified as $T_{EX}^{int\text{-}P6}$ cells because they manifested signatures of both intermediately exhausted (T_{EX}^{int}) and T_{EX} cells, and flowed directly into cluster P8 cells (**Fig 2A-D**), which were designated $T_{EX}^{term\text{-}P8}$ cells because they were strongly enriched with the T_{EX}^{term} cell signature (**Fig 1D,** fig S1M) and highly expressed *Tox, Pdcd1* and *Lag3* (**Fig 1E and G**). Thus, T_{EX}^{int} cells are related to classical MP cells, but exhibit altered transition probabilities into

terminal states and fail to establish the T_{CIRC} lineage during $LCMV_{Cl13}$ infection (**Fig 2B-D** and fig S2C).

Trajectory 4 (T4: P2> P7) defined formation of precursors of T_{SCM}-like cells during acute (T_{SCM}^{P7}) or chronic ($T_{EX}^{prog1\text{-}P7}$) LCMV infection (**Fig 2C**). T_{SCM}^{P7} cells from $LCMV_{Arm}$- infected hosts were positively enriched with T_{SCM} and T_{CM} cell signatures, and were strongly connected to pT_{EM}^{P4} and EE^{P9} cells (**Fig 1D** and fig S1J-K). Both T_{SCM}^{P7} and pT_{EM}^{P4} cells exhibited strong *Tbx21* RNA velocity, suggesting they both manifest transition potential into *Tbx21* expressing states. Consistent with this, cells in both clusters exhibited strong transition probabilities into EE^{P9} cells, indicating that T_{SCM} precursors during $LCMV_{Arm}$ infection are poised with T_{EFF} cell potential. In contrast, although $T_{EX}^{prog1\text{-}P7}$ cells from $LCMV_{Cl13}$-infected hosts were positively enriched with the T_{SCM} signature, they exhibited reduced *Tbx21* RNA velocity (Table S5), and lacked transition probability into EE^{P9} cells (**Fig 2B**). In addition, $T_{EX}^{prog1\text{-}P7}$ cells appeared to derive from $EMP^{P2\text{-}Cl13}$ cells initially, but their strong accumulation by day 8 pi correlated with retrograde transition potentials from all downstream T_{EX} cell clusters except $T_{EX}^{prog2\text{-}P3}$ cells, whereas during $LCMV_{Arm}$ infection, T_{SCM}^{P7} cells derived predominantly from EMP^{P2} cells (**Fig 2B-D**). Moreover, $T_{EX}^{prog1\text{-}P7}$ cells more strongly induced *Tcf7* (**Fig 2D**), highly expressed *Lag3*, *Pdcd1*, *Tox*, *Tox2* and *Bcl6*, were positively enriched with the specific T_{EX} cell signature and were enriched with terminal states (**Fig 1D** and figs S1H, M and S2C). Thus, the precursors of T_{SCM} cell states during acute and chronic infections have different origins and distinct developmental potentials.

IL-2R-dependent transcription establishes EMP cells and transcriptional heterogeneity

The most dynamic genes drive the RNA velocity vector field [11,12]. Those encoding TFs, CRFs, and surface receptors (SRs) during transitions between clusters were identified as potential drivers that compose the antiviral CD8 T cell architecture (**Fig 2C** and fig S2D-E, Table S4). Strong *Il2ra* dynamics and its transiently high expression in EMP^{P2} cells indicated IL-2R-dependent signals might promote multilineage potential (**Fig 3A and B**). To define the role of *Il2ra* functionally, single cell trajectories were inferred after scRNA-seq analysis of wildtype and *Il2ra*-deficient P14 (P14 *Il2ra*$^{-/-}$) CD8 T cells 6 days after $LCMV_{Arm}$ infection (**Fig 3C and D,** as described fig S1B). This second analysis confirmed the original lineage architecture during $LCMV_{Arm}$ infection (**Fig 3C** and refer to **Fig 1A**). Louvain clusters from the *Il2ra* analysis (designated "Exp-*Il2ra*") closely matched most clusters identified on days 5 and 8 pi in the original analyses (**Fig 3D** and refer to **Fig 1**). PAGA-inferred connectivity and RNA velocities implied analogous intracluster transition probabilities (**Fig 3C and E**), and analogous single cell

behavior of wildtype P14 cells in both the original and Exp-*Il2ra* analyses emphasized that the inferred lineage architecture is biologically robust.

P14 *Il2ra*$^{-/-}$ CD8 T cells distributed within the trajectories aberrantly compared to WT P14 cells. P14 *Il2ra*$^{-/-}$ cells almost entirely composed cluster Exp-*Il2ra*^{P1} compared to wildtype P14 cells indicating they arrested before transition into EMP^{P2} cells (Exp-*Il2ra*^{P2}) (**Fig 3F** and fig S3E, p-value 1.35 x 10^{-27}). Exp-*Il2ra*^{P1} cells were activated (data not shown), but lacked RNA velocity into future states indicating they were terminal (fig S3D). Strong differenital expression between Exp-*Il2ra*^{P1} and Exp-*Il2ra*^{P2} cells confirmed *Il2ra* was essential for transition into EMP^{P2} cells (**Fig 3I**). In addition, differential expression between wildtype and *Il2ra*$^{-/-}$ P14 cells in cluster Exp-*Il2ra*^{P2} confirmed IL-2 signaling was required for EMP cell formation (fig S3G). This required IL-2R-dependent transcription, because genes whose nascent RNA expression required *Il2ra* for upregulation after TCR stimulation (WT^{48h} > WT^{naive}, padj < 0.05) were positively enriched with those upregulated as Exp-*Il2ra*^{P1} transitioned into Exp-*Il2ra*^{P2} cells (fig S3K, NES = 1.26, pvalue = 0.04), whereas genes that required *Il2ra* for downregulation (WT^{48h} > *Il2ra*$^{-/-,48h}$, padj < 0.05) were positively enriched with those downregulated in this transition (fig S3K, NES = -1.16, pvalue = 0.05). Thus, IL-2R-dependent transcription in vivo is essential for gene expression that drives activated naive cells to become EMP^{P2} cells.

Development beyond the EMP^{P2} cell state also required IL-2R signals. The transition probabilities of P14 *Il2ra*$^{-/-}$ cells into other clusters were substantially altered (**Fig 3E**). *Il2ra*$^{-/-}$ cells in cluster Exp-*Il2ra*^{P2} lacked transition potential (**Fig 3E**). Those from cluster Exp-*Il2ra*^{P4} did not manifest velocity into Exp-*Il2ra*^{P5} (T_{RM}) cells, and those in Exp-*Il2ra*^{P5} were vectored backward into cluster Exp-*Il2ra*^{P2} (**Fig 3E**). Consistent with this, P14 *Il2ra*$^{-/-}$ cells were depleted from cluster Exp-*Il2ra*^{P5} cells (pT_{RM}^{P3} analog) (**Fig 3F** and fig S3E, p-value 0.057), and those that did accumulate in that cluster were negatively enriched with the T_{RM}-signature compared to wildtype P14 Exp-*Il2ra*^{P5} cells (fig S3F). Thus, *Il2ra*$^{-/-}$ P14 cells that bypassed the initial developmental block inefficiently formed putative T_{RM} precursors. In addition, *Il2ra*$^{-/-}$ cells in cluster Exp-*Il2ra*^{P4} manifested retrograde vectors into Exp-*Il2ra*^{P3} (T_{SCM}) cells, unlike wildtype P14 cells (**Fig 3E**), which correlated with increased P14 *Il2ra*$^{-/-}$ cell representation in cluster Exp-*Il2ra*^{P3} (T_{SCM}^{P7} analogs) (**Fig 3F** and fig S3E, p-value 0.00042). Furthermore, *Il2ra*-deficient cells inefficiently repressed T_{SCM} signature genes after TCR stimulation (fig S3J). These results demonstrate that IL-2R signals promote divergent transcription in EMP cells that establishes trajectories into branching T_{MEM} cell lineages.

The bias of EMP-like *Il2ra*$^{-/-}$ cells toward T_{SCM}-like states and their reduced contribution to other T_{MEM} precursor states during acute infection prompted examining IL-2R-regulated genes in EMP^{P2} cells from $LCMV_{Cl13}$-infected mice. EMP^{P2} cells during $LCMV_{Cl13}$ infection less highly expressed IL-2R-Stat5 induced genes (*Batf3*, *Ccr5*, *Gzmb*, *Chd7* see below) that promote formation of protective T_{EFF} and T_{MEM} cells (fig. S3L). The IL-2R-repressed gene signature was enriched with genes whose RNA velocity was greater in EMP^{P2} cells from $LCMV_{Cl13}$-infected mice (fig S3M). Moreover, in $LCMV_{Cl13}$ infection, $T_{EX}^{prog1\text{-}P7}$ cells upregulated the IL-2R-repressed gene signature compared to EMP^{P2} cells; whereas this signature was not upregulated as EMP^{P2} transitioned to T_{SCM}^{P7} cells during $LCMV_{Arm}$ infection (fig S3M). Thus, regulation of IL-2R-dependent genes appear to bias the future states of EMP^{P2} cells during acute and chronic infection.

Differential utilization of CRFs establishes antiviral CD8 T cell heterogeneity

Chromatin structure in naive cells is remodeled during initial TCR and IL-2 stimulation and becomes stably differential in distinct T_{EFF}, T_{MEM} and T_{EX} cell subsets[38]. To identify chromatin regulatory factors (CRFs) that might control diverging transcriptional programs during T_{EFF} and T_{MEM} cell formation we screened a library of retrovirally delivered microRNA-embeded short hairpin RNAs (shRNAmirs) targeting nearly all murine CRF genes using a pooled approach in P14 CD8 T cells responding to $LCMV_{Arm}$ infection (fig S4A and B, **Fig 5A** and Table S7)[14]. Candidate CRFs were identified by sequencing DNA libraries amplified from integrated shRNAmir proviral sequences in FACS-purified CD8 T cell subsets and quantifying differential shRNAmir representation (RNAi-induced effects) (**fig S4B**, https://github.com/Yolanda-HT/HSAP). Genes with top RNAi-effects (25th percentiles) affecting in vivo P14 cell accumulation (input vs output), maturation of KLRG1^{lo} CD127^{lo} cells into all other phenotypes (EE vs other), and the balance between TE and MP cells (KLRG1^{hi}CD127^{lo} vs KLRG1^{lo}CD127^{hi}) were selected as potential candidates (**Fig 4B-E** and Table S7). Individual follow-up experiments confirmed specific phenotypes of several top candidates that were previously unknown, including *Prmt5*, *Carm1*, *Taf1, Mll1* (manuscript in preparation) and multiple genes encoding factors in Brg1-associated factor (BAF, mammalian SWI/SNF)[39] and chromodomian helicase and DNA-binding (Chd) nucleosome remodeling complexes (**Fig 4B-D**). Thus, concerted activity of many CRFs differentially control formation of classically defined populations defined by KLRG1 and CD127 expression during acute viral infection.

We pursued *Smarca4* (Brg1) and *Chd7* (Chd7), which encode ATPases of nucleosome remodeling complexes that have essential roles in multipotent neural crest cells[40], human development and immune system function[41-43], and thymic T cell development[44]. Their strong phenotype in the screen and their dynamics in the single-cell trajectory, both implied they could have essential functions in EMP cells. Depletion of either factor alone impaired formation of KLRG1^{hi} cells on day 5 pi, and increased the fractions of KLRG1^{lo} CD127^{lo} EE-like cells by day 8 pi, which suggested both factors might promote divergence into TE, DP or MP populations[4] (**Fig 5B**). In addition, RNAi-effects in the primary screen indicated *Smarca4* and at least 4 additional BAF subunits (*Arid1a, Smarcc1*, *Smarce1*) were selectively required for TE cell formation (**Fig 4C**). Four other BAF-subunits (*Actl6b*, *Smarcc2*, *Smarcd1*, and *Smarcd3*) were preferentially required for MP formation (**Fig 4C**). Thus, distinct BAF-complex assemblies might differentially bias TE and MP cell development[39]. To further confirm the role of Chd7, engineered *Chd7* alleles[42] were conditionally disrupted in mice using transgenic Cre expression in post-thymic T cells (referred to as *Chd7*$^{fl/fl}$ dLck-Cre sfYFP). Similar to RNAi, *Chd7* gene-disruption impaired the frequency of KLRG1^{hi} cells 5 days after LCMV$_{Arm}$ infection (fig S4C), and reduced formation of TE cells while increasing frequencies of EE- and MP-like cells 8 days after infection, without strongly affecting overall T_{EFF} cell numbers (**Fig 4E**). Thus, both BAF and Chd7 complexes are essential for the early phenotypic heterogeneity of T_{EFF} cells during acute infection.

Smarca4 and Chd7 are required for transcriptional heterogeneity of EMP cells

Distinct RNA dynamics of *Smarca4* and *Chd7* in the trajectory implied sequential early functions during initial lineage divergence. Mature *Smarca4* mRNA expression was greatest in EMP^{P2} cells, before *Chd7* RNA velocity increased in EMP^{P2}, DP^{P5} and MP^{P6} cells (**Fig 4G**). Mature *Chd7* transcripts were upregulated in EE^{P9} and TE^{P10} cells (**Fig 4G**) and in bulk KLRG1^{hi} cells on day 5 of LCMV$_{Arm}$ infection[25]. *Chd7* transcription required *Il2ra* during TCR stimulation (**Fig 3K**). These data are consistent with Smarca4 and Chd7 functioning in an IL-2R-dependent transctiptional network. To examine this possibility, P14 cells depleted of either *Smarca4* or *Chd7* were analyzed by RNA-seq 5 days after LCMV$_{Arm}$ infection. *Smarca4* was required to downregulate genes that are repressed by IL-2R (day 6), and to repress both the T_{SCM} and *Tcf7*-promoted gene expression signatures [21,25,36,45-47] (**Fig 4H**), whereas *Chd7* was required for activation of genes that require IL-2R for expression at later times (day 10)[46] (**Fig 4H**). In addition, both factors were necessary for promoting gene expression activated by the TF Runx3

and repressing gene expression promoted by the TF Tox (**Fig 4H**). Thus, IL-2R-dependent transcriptional heterogeneity that develops in EMP cells early during acute infection required both Smarca4 and Chd7. In addition, the distinct requirements of each factor indicated they differentially promote transcription that drives divergence between T1 (pT_{EM}), T2 (MP) and T4 (pT_{SCM}) trajectories.

Smarca4 and Chd7 stabilized transcription that drives formation of the T1 trajectory. *Smarca4*-depleted P14 cells on day 5 pi expressed significantly less *Bhlhe40, Chd7, Gzma, Med12l, Runx3*, *Tbx21* and *Zeb2* (**Fig 4I**, *left*); *Chd7*-depleted cells expressed significantly less *Gzma, Il12rb2*, *Il18rap*, *Med12l* and *Zeb2*, while expression of *Bhlhe40, Batf3,* and *Tbx21* trended lower (**Fig 4I**, *right*). In P14 cells depleted of Chd7, T-bet protein expression and TE cell formation was more strongly impaired during LM_{GP33} infection than during $LCMV_{Arm}$ infection, consistent with grossly impaired *Il12rb2* expression (**Fig 4I**, *right*), and the increased IL-12 concentrations during LM infection compared to LCMV infection[30,48] (**Fig 6J-K** and data not shown). Complementation of Smarca4 or Chd7 depleted P14 cells with retrovirally expressed Tbx21 restored the normal pattern of TE and MP in each case (fig 4D and E), indicating they each promote T_{EM} and TE differentiation by ensuring *Tbx21* expression. However, enforced T-bet expression in the absence of *Chd7* did not resuce defective Gzmb expression, indicating that Chd7 is broadly required for cytolytic effector cell programming (data not shown). Because altered gene expression in the absence of Smarca4 and Chd7 is manifest on day 5 pi, prior to EE^{P9} cell formation, these results demonstrate that both CRFs are necessary to establish T_{EFF} gene expression prior to when cells with these phenotypes manifest.

Chd7 is essential for T_{MEM} cell lineage branching

P14 cells depleted of Chd7 during $LCMV_{Arm}$ infection were positively enriched with gene expression signatures of DP^{P5} and MP^{P6} cells on day 5 pi, which correlated with increased frequencies of EE and MP-phenotype cells in *Chd7*$^{fl/fl}$ dLck-Cre sfYFP mice 8 days after $LCMV_{Arm}$ infection (**Fig 4F**). Thus, cells lacking Chd7 appeared to arrest at the point where EMP^{P2} cells undergo branching into MP^{P6} and EE^{P9} cells, well before maturation of TE^{P10} cells, which brought into question whether they correctly stabilized the specific gene expression programs related to each of these flow cytometry phenotypes. To examine this directly, LCMV-specific CD8 T cells from *Chd7*$^{+/+}$ and *Chd7*$^{fl/fl}$ dLck-Cre^{+} sfYFP^{+} mice that exhibited MP, EE, DP and TE cell phenotypes 8 days after $LCMV_{Arm}$ infection were FACS-purified and analyzed using RNA-seq. Multidimensional scaling showed these populations from wildtype mice were

separated from each other, whereas those from *Chd7*-deficient mice grouped (**Fig 4L**), indicating that gene expression states which diverged in wildtype T_{EFF} subsets did not strongly diverge in the *Chd7*-deficient populations. Consistent with this interpretation, pairwise analysis showed that compared to wildtype cells, *Chd7*-deficient TE-phenotypic cells less strongly expressed genes encoding factors characteristic of TE cells (*Zeb2*, *Med12l*, *Il18rap, Il12rb2*), and *Chd7*-deficient EE- or MP-phenotypic cells less highly expressed genes that promote MP cell development and formation of long-lived T_{MEM} cells (*Tcf7*, *Id3*, *Tox,* and *Ccr7*) (**Fig 4M**). These results demonstrate that Chd7 is necessary to stabilize divergent transcriptional programs that differentiates circulating T_{MEM} lineage branches and maturing cell states, and promotes terminal T_{EFF} maturation.

Discussion

Our study resolves the initial stages of antiviral CD8 T cell ontogenesis. The architecture indicates naive cells differentiate into common EMP cells which diverge along distinct trajectories that develop gene expression states within the first week of acute or chronic viral infection that match all major T_{EFF}, T_{MEM} and T_{EX} cell populations found at later times. Additional developmental paths to cells that were not sampled in this analysis could exist (e.g., cells from other tissues)[49,50]. Variable RNA velocities that develop in EMP cells indicates that diverging transcription initiates the branching trajectories before strong differential expression of lineage-specific regulators is established. IL-2R signals were required for EMP cell development and their transcriptional heterogeneity, and altered IL-2R-dependent transcription in EMP cells during chronic infection correlated with development of T_{EX} cell states. Because *Il2ra* is regulated by IL-2 stimulation and was dynamic during EMP cell formation and divergence, these results suggest variable IL-2 stimulation contributes to initial lineage bias of cells in the population.

We provide evidence that diverse T_{MEM} cell types most likely arise from divergent precursor cell states derived from distinct lineages early in the response. The nucleosome remodeler Smarca4 was necessary for gene expression that initially separates T_{SCM} and T_{EM} precursor states; Chd7 functioned later to mature classical MP cells; and, both factors cooperatively promoted TE cell differentiation. These results provide evidence that diverging transcriptional states in EMP cells are stabilized by specific CRFs, and implies that chromatin remodeling reinforces initial lineage biases and establishes the branching architecture. Thus, differential chromatin remodeling in divergent T_{MEM} cell precursor lineages might explain the

preferential interconversion potentials of distinct mature T_{MEM} cell populations[1,51,52], and resistance of T_{EX} cells to chromatin-level reprogramming[53,54]. The developmental architecture, stepwise transcriptional dynamics and CRF atlas described here suggests many factors with spatiotemporally resolved functions and might suggest strategies for engineering T_{MEM} CD8 T cell formation.

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Figure 1. Naive CD8 T cells differentiate along a linear path into common effector and memory progenitor (EMP) cells.

(A) PAGA initialized force-directed (FA) embedding based on individual cell gene expression profile, each dot represent one cell. PAGA connectivity analysis was performed on indicated clusters P0-P10. Clusters were extracted by Louvain method based on neighborhood graph. Neighborhood graph was calculated with UMAP algorithm.

(B) Top: PAGA connectivity graph, each node represent one cluster, node sizes represent relative cell number of cluster, edge widths represent relative PAGA connectivity score. Bottom: heatmap of PAGA connectivity score between clusters.

(C) PAGA initialized FA embedding, coloring cells based on origin: day5 /day8, naive (grey) / $LCMV_{Arm}$ (purple) / $LCMV_{Cl13}$ (orange).

(D) GSEA of selected signatures for Louvain clusters, separating $LCMV_{Arm}$ / $LCMV_{Cl13}$. GSEA analysis performed on mean of normalized gene expression per cluster. NES represented by color, -log10(padj) represented by dot size. Labels of clusters inferred by GSEA result. (See also fig S1 J-M)

(E) Mean of scaled expression, showing top signature genes of Louvain clusters. Signature genes were selected based on multiple differential analysis (Wilcoxon rank sum test, t-test, t-test over estimated variances) of cells within each cluster v.s. all others, with adjusted p-adj cutoff 0.05 (intersection of all tests) and absolute log2 fold change cutoff 1. Chromatin remodeling factors, transcription factors and surface proteins were selected from the genes that meet the statistical cutoffs for creating the signature gene lists. (See also Table S1). Top 10 signature genes ranked by t-test scores are represented in heatmap.

(F) Expression of cell surface markers by CTV determined by flow cytometry. 50,000 P14 CD8 T cells were transferred per recipient mice which were given $LCMV_{Arm}$. Naive cells (grey) and cells from day3 (blue) / day4 (purple) post infection are represented in plots.

(G) Raw expression (logrithmized and normalized) of selected genes.

Figure 2. Disparate RNA velocities develop in individual EMP cells and initiate a branched manifold that establishes T_{MEM} and T_{EX} cell diversity.

(A) Stream plot of velocity embedded on PAGA initialized single cell projection, separating $LCMV_{Arm}$ / $LCMV_{Cl13}$ (including naive cells in both conditions).

(B) Transition probability heatmap between clusters estimated by scVelo, separating $LCMV_{Arm}$ / $LCMV_{Cl13}$. Color represent transition probability from row cluster to column cluster.

(C) Inferred $LCMV_{Arm}$ / $LCMV_{Cl13}$ developmental rajectory by PAGA connectivity analysis, scVelo transition probability, pseudo-time (see Table S1) and real time.

(D) Spliced transcript abundance (Ms) of representative genes for each of the four inferred trajectory in $LCMV_{Arm}$ / $LCMV_{Cl13}$.

(E) Single cell velocity of selected driver genes, separating $LCMV_{Arm}$ / $LCMV_{Cl13}$. Potential driver genes were identified by combining top likelihood genes from analysis of all cells or multiple pairs of clusters with transitioning potential. Transcription factors, chromatin remodelers and surface receptors were selected within the identified likelihood genes (See Table S3).

Figure 3: IL-2R-dependent transcription establishes EMP cells and transcriptional heterogeneity.

(A) Heatmap of spliced transcript abundance of EMP^{P2} signature driver genes. EMP^{P2} common driver genes were defined as intersection between driver genes with likelihood > 0.25 in: all cells, P2-P4, P2-P5 and P2-P7 (Table S3) P2 signature driver genes were defined as intersection between P2 signature genes (described in **Fig 1E** and Table S1) and EMP^{P2} common driver genes.

(B) Single cell normalized Il2ra transcript abundance, spliced (Ms) / unspliced (Mu).

(C) PAGA initialized FA embedding based on single cell cell gene expression profile for Exp-*Il2ra*. PAGA connectivity analysis was performed on indicated clusters. Clusters were extracted from UMAP neighbor graph by Louvain method.

(D) Correlation of clusters between acute v.s. chronic single cell experiment and Exp-*Il2ra*. Correlation represented by mean UMAP nearest neighbor graph scores between clusters from two experiments. UMAP projection of single cells from two experiments generated by Harmony integrated normalized count matrix from both experiment.

(E) Transition probability between clusters in activated WT and *Il2ra*$^{-/-}$ cells. Left: all single cells from WT / *Il2ra*$^{-/-}$ samples in PAGA projection. Right: transition probability heatmap.

(F) Percentage distribution in Louvain clusters for each cell type. Total percentage in all Louvain clusters add up to 100% for each cell type.

(G) Signature gene heatmap for clusters (calculated for all cells, including naive, WT and *Il2ra*$^{-/-}$). Method as described in **Fig 1E**.

(H) Heatmap of log2 fold change of WT versus *Il2ra*$^{-/-}$ for selected differential genes. Genes were selected from cluster signature gene list (described in **Fig 2G**). The genes with minimum expression of 0.0015 in at least one cluster and with absolute log2fc >= 2 in at least one cluster (WT versus *Il2ra*$^{-/-}$) were used.

(I) Differential analysis of transcript abundance comparing Exp-Il2ra^{P1} and Exp-Il2ra^{P2}. X-axis represents log2 fold change, and the y-axis represents -log10(pvalue). The horizontal line indicates pval = 0.05. The vertical lines indicates absolute log2fc = 1. T_{RM} and T_{SCM} signature genes are highlighted in yellow and green respectively.

(J) Number of differential nascent transcripts (between WT and *Il2ra*$^{-/-}$) at different time points post activation. Differential nascent transcript is determined by DESeq2 (padj < 0.05) .

(K) Differential analysis volcano plot of WT versus *Il2ra*$^{-/-}$ at 48 hours post activation. The x-axis represents log2 fold change, and y-axis represents -log10(pvalue). The horizontal line indicates pval = 0.05. The vertical lines indicates absolute log2fc = 1. T_{RM} and T_{SCM} signature genes are highlighted in yellow and green respectively.

Figure 4: Differential utilization of CRFs establishes antiviral CD8 T cell heterogeneity.

(A) Pooled RNAi screen and analysis simplified schematic (detailed schematic in fig S4A-B).

(B-D) Ranked lists of adjusted RNA - effects for input versus output, MP versus TE, and EE versus others. Red / blue highlight genes that are top / bottom quarter in effect ranking.

(E-F) Experiment characterizing endogeneous CD8^{+} T cells response to LCMV$_{Arm}$ infection (day 8 pi), in mice of Chd7$^{fl/fl}$, Chd7$^{+/fl}$, Chd7$^{+/+}$ genotypes. (E) Total activated CD8 T cell number in spleens. (F) Representative flow cytometry plots showing CD127 and KLRG1 staining, and summarized percentages of cells in each category.

(G) Smarca4 and Chd7 velocity and spliced transcript abundance (Ms) in single cell projection, separating LCMV$_{Arm}$ / LCMV$_{Cl13}$.

(H) GSEA of sh*Chd7* versus sh*Cd4* (control) and sh*Brg1* versus sh*Cd19* (control). Signatures: IL-2 regulated signatures / key CD8 transcription factor (TF) regulated signatures / CD8 phenotype signatures / Best et. al. longitudinal expression dynamic gene signatures.

(I) Differential expression of selected genes. Heatmap showing log2 fold change of gene expression comparing sh*Chd7* versus sh*Cd4* (control) or sh*Brg1* versus sh*Cd19* (control).

(J-K) Comparison of T-bet expression and phenotype between sh*Chd7* versus sh*Cd4* (control) during LCMV$_{Arm}$ or Listeria infection. Top: representative and summarized T-bet MFI. Bottom: representative flow cytometry plot showing CD127 and KLRG1 staining.

(L) Multidimensional scaling plot showing similarity / dis-similarity of Chd7$^{fl/fl}$ versus Chd7$^{+/+}$ cells in different stages: naive / 48h post activation / day8 sorted subsets.

(M) Differential expression of selected genes. Heatmap showing log2 fold change of gene expression between Chd7$^{fl/fl}$ versus Chd7$^{+/+}$ in day8 sorted subsets (TE, MP, EE).

Supplementary Information:

- Figures S1-S4
- Tables S1-S7

Figure S1. Unsupervised approach to define early developmental clusters of antiviral CD8 T cells during actue and chronic viral infections at single cell level.

(A) Schematic of acute v.s. chronic single cell experiment. Donor naive P14 Thy1.1 $CD8^+$ T cells from were isolated and transferred into 4 groups of WT recipient mice at day -1. $LCMV_{Arm}$ and $LCMV_{Cl13}$ infections were given at either day 0 or day 3 for each group. P14 Thy1.1 donor $CD8^+$ T cells and GP33 $Tetramer^+$ endogenous LCMV responding $CD8^+$ cells (Tet) were isolated at the same day for each group. Cells of each different origin were hash-tagged and mixed for scRNAseq in the same batch.

(B) Schematic of bioinformatic analysis pipeline of acute v.s. chronic single cell experiment. 10x outputs were converted to fastq files and counted for transcript abundance with Cellranger 3.0. Basic quality filter and normalization was performed with Scanpy. Biolegend hashtags were demultiplexed with custom python script. Outliers were detected with DBSCAN (scikit-learn). Cells from each condition were resampled to 750 – 1250 cells condition. Dimentionality reduction (PCA, UMAP, Force Atlas), clustering (Louvain) and cluster connectivity analysis (PAGA) were performed with Scanpy functions. The count matrix was also processed with Velocyto to separate spliced and unspliced transcripts for further velocity associated analysis with scVelo.

(C) Stacked bar chart showing composition by different cell type for each Louvain cluster P0-P10. For each cluster, total percentages of all cell types add up to 100%.

(D) Heatmap showing percentage distribution in Louvain clusters for each cell type. For each cell type, total percentage in all Louvain clusters add up to 100%.

(E) Heatmap represents similarity between cell types based on distribution in Louvain clusters estimated by bhattacharyya coefficient.

(F) Chi-square analysis of distribution of day 5 $LCMV_{Arm}$ / $LCMV_{Cl13}$ P14 $CD8^+$ cell distribution in / out cluster P2.

(G) Violin plots of raw expression (logrithmized and normalized) per cluster for selected phenotype marker genes.

(H) See **Fig 1E** description: mean of scaled expression heatmap, top 10 genes ranked by t-test score plotted. Left: all cells; middle and right: separating $LCMV_{Arm}$ / $LCMV_{Cl13}$.

(I) Phenotype correlation between CX3CR1 - CD27 and CD25 - TCF-1 of transferred P14 $CD8^+$ cell day 4 post LCMV infection (50k cells transferred).

(J) (K) (L) (M) GSEA of selected signatures for Louvain clusters, separating $LCMV_{Arm}$ / $LCMV_{Cl13}$. GSEA analysis performed on mean of normalized gene expression per cluster. Colors of dots represent NES, and sizes of dots represent -log10(padj) of signature enrichment. Best et. el. signatures annotated with longitudinal expression dynamics of genes in the clusters and grouped based on expression dynamics (plot from original publication).

Figure S2. Single cell RNA velocity and root / terminal score indicates P2 & P5 are developmental roots.

(A) Single cell velocity vectors for each cluster.

(B)(C) Root / terminal score analysis, separating $LCMV_{Arm}$ / $LCMV_{Cl13}$ (not including naive cells). Top: Root / terminal score of single cells, overlayed with grid velocity. Bottom: average root / terminal per cluster.

(D)(E) Single cell spliced transcript abundance / unspliced transcript abundance for selected driver genes, Separating $LCMV_{Arm}$ / $LCMV_{Cl13}$.

(F) GSEA analysis of P14 $Runx3^{-/-}$ versus WT CD8 T cell gene expression, at day 5 / day 8 p.i. LCMV infection. Positive NES scores indicate signature genes more highly expressed in $Runx3^{-/-}$ comparing with WT; negative NES scores indicate signature genes more highly expressed in WT comparing with $Runx3^{-/-}$.

Figure S3. IL-2 signaling contribute to EMP formation via transcription regulation

(A) Schematic of Exp-*Il2ra*: *Il2ra*$^{-/-}$ and WT P14 $CD8^+$ T cell cogenetic transfer and single cell experiment.

(B) Left: PAGA connectivity graph, each node represent one cluster, node sizes represent relative cell number of cluster, edge widths represent relative PAGA connectivity score. Middle: heatmap of PAGA connectivity score between clusters. Right: Stacked bar chart showing composition by different cell type for each Louvain clusters. For each cluster, total percentages of all cell types add up to 100%.

(C) Exp-*Il2ra* PAGA initialized FA embedding, highlighting cells of different origin.

(D) Terminal score of *Il2ra*$^{-/-}$ single cells in Exp-*Il2ra*.

(E) Chi-square analysis of cell type distribution in Exp-*Il2ra*. Top: Cell number distribution in activated clusters in Exp-$Il2ra^{P1}$ or outside of Exp-$Il2ra^{P1}$ for WT and I *Il2ra*$^{-/-}$. Bottom: Cell number distribution in activated clusters except for P1, comparing WT and $Il2ra^{-/-}$.

(F) GSEA of T_{RM} gene signature in Exp-Il2ra^{P5} WT versus *Il2ra*$^{-/-}$.

(G) Differential expression volcano plot of WT and *Il2ra*$^{-/-}$ in Exp-Il2ra^{P2}. The x-axis represents log2 fold change, and y-axis represents -log10(pvalue). The horizontal line indicates pval = 0.05. The vertical lines indicates absolute log2fc = 1. TE signature genes marked in green.

(H) Schematic of *Il2ra*$^{-/-}$ and WT P14 CD8^{+} T cell *in vitro* activation and nascent RNA-seq experiment.

(I) See **Fig 3H**. Dark red and dark blue area represent number of genes genes that are expressed higher in WT than *Il2ra*$^{-/-}$ which are upregulated post activation (each time point versus naive), or genes expressed lower in WT than *Il2ra*$^{-/-}$ which are downregulated post activation.

(J) GSEA of T_{SCM} signature indicates that *Il2ra*$^{-/-}$ express T_{SCM} signature genes at higher level than WT at 48h post activation.

(K) GSEA enrichment of IL-2 promoted / repressed genes in Exp-Il2ra^{P3}(EMP) versus Exp-Il2ra^{P1} (Arrested) differential gene list (signature genes selected from DEseq2 analysis of 48h post activation, WT versus *Il2ra*$^{-/-}$ with cutoffs padj < 0.05, absolute log2 fold change > 1.2).

(L) Venn diagrams showing overlap between P2 $LCMV_{Arm}$ and $LCMV_{Cl13}$ differential genes (padj < 0.05) and IL2-Stat5 promoted / repressed genes. Pie charts showing the percentage of Stat3 bound genes in the intersection of $LCMV_{Arm}$ and $LCMV_{Cl13}$ differential genes and IL2-Stat5 promoted / repressed genes. IL2-Stat5 promoted genes: nascent transcript abundance WT > *Il2ra*$^{-/-}$ and WT 48h > WT 6h (DESeq2, pvalue <= 0.05), intersected with genes that are bound by Stat5 [55]. IL2-Stat5 repressed genes: nascent transcript abundance WT < *Il2ra*$^{-/-}$ and WT 48h < WT 6h (DESeq2, pvalue <= 0.05), intersected with genes that are bound by Stat5.

(M) GSEA enrichment of IL2 repressed signature between $LCMV_{Arm}$ and $LCMV_{Cl13}$ differential velocity genes in EMP^{P2}.

(N) GSEA enrichment of 48h IL2 repressed signature (described in fig S3K) in T_{SCM}^{P7} versus EMP^{P2}, separating $LCMV_{Arm}$ and $LCMV_{Cl13}$.

(O) Visualization of ChIP-seq, ATAC-seq and nascent RNA-seq tracks at *Il6st* and *Bcl6* regions. Asterisks represent significant peaks (MACS2, q-value < 0.01).

Figure S4. Identification and functional analysis of CRFs that is required for CD8^{+} T cell lineage formation

(A) Experimental setup of CRF RNAi screen (See Methods).

(B) Bioinformatic analysis pipeline of CRF RNAi screen ().

(C) Flow cytometry at day5 pi showing KLRG1^{hi} population in Chd7$^{fl/fl}$ is reduced comparing with Chd7$^{+/+}$.

(D)(E) Flow cytometry of transduced and transferred P14 CD8 T cells post LCMV$_{Arm}$ infection. CD8 T cells were simultaneously transduced with a combination of retroviral vectors containing sh*Cd4* (Ametrine, control) and empty vector (GFP, control), or a combination of retroviral vectors containing sh*Smarca4* / sh*Chd7* (Ametrine) and T-bet cDNA (GFP). Phenotypes of RNAi and T-bet overexpression were accessed by flow cytometry at day8 pi.

(F)(G)(H) Ranked lists of adjusted RNA - effects for input versus output, MP versus TE, and EE versus others (as described in fig 5B-D), annotating genes from major CRF families.

(I) Six major clusters of CRFs extracted from hierarchical clustering of CRFs based on RNAi effect scores of 3 categories (MP v.s. TE, EE v.s. Other, Input v.s. Output) from screen.

(J)(K) Flow cytometry of splenic CD8^{+} T cell phenotype day8 pi LCMV$_{Arm}$, comparing transferred P14 CD8 $^{+}$ T cells transduced with sh*Cd4*(control) and multiple shRNAs against Carm1 and Prmt5.

(L) Flow cytometry of peripheral blood CD8^{+} T cell phenotype day7 and day12 pi LCMV$_{Arm}$, comparing congenially transferred P14 CD8 $^{+}$ T cells transduced with sh*Cd19*(control) and sh*Taf1*.

Table S1.

Acute versus Chronic infection (LCMV$_{Arm}$ / LCMV$_{Cl13}$) single cell experiment - clusters and cluster signature genes

Table S2.

GSEA signatures

Table S3.

Acute versus Chronic infection (LCMV$_{Arm}$ / LCMV$_{Cl13}$) single cell experiment - differential expression of genes between LCMV$_{Arm}$ and LCMV$_{Cl13}$

Table S4.

Acute versus Chronic infection (LCMV$_{Arm}$ / LCMV$_{Cl13}$) single cell experiment - scVelo likelihood genes

Table S5.

Acute versus Chronic infection ($LCMV_{Arm}$ / $LCMV_{Cl13}$) single cell experiment - spliced / unspliced transcript abundance and velocity of likelihood genes

Table S6.

Exp-*Il2ra* - clusters and cluster signature genes

Table S7.

RNAi screen of chromatin remodelers: shRNA sequences; scores of each gene in different comparisons

Materials and Methods

Mice

Wildtype 6-8 week old C57BL/6J mice were used as recipients for adoptive transfer experiments and were purchased from the Jackson Laboratory. P14 Thy1.1^{+} mice were a gift from Dr. Rafi Ahmed (Emory University). P14 Thy1.1^{+} Thy1.2^{+} mice were generated by crossing P14 Thy1.1^{+} mice with wildtype C57BL/6J mice. P14 Thy1.1^{+} *Il2ra*$^{-/-}$ mice were generated by crossing *Il2ra*$^{-/-}$ mice (purchased from Jackson Laboratory) with P14 Thy1.1^{+} mice. *Chd7*$^{fl/fl}$ mice were a gift from Dr. Donna M. Martin (University of Michigan) [42], and were crossed with Rosa26-EYFP and dLck-Cre (maintained heterozygous) transgenic mice. All mice were maintained in specific-pathogen free facilities and used according to protocols approved by the Institutional Animal Care and Use Committee of TSRI-FL.

T cell activation, adoptive transfer and infections

Naive P14 CD8^{+} T cells from wildtype mice were isolated by negative selection (EasySep™, Stemcell Technologies). Naive P14 Thy1.1^{+} *Il2ra*$^{-/-}$ cells were isolated from 4-5 week old mice by depleting CD44^{hi} cells (Biolegend biotin anti-mouse/human CD44, clone IM7, 2µl per spleen). For *Il2ra*$^{-/-}$ single cell experiment, cells were further sorted for CD44^{lo} with FACS. Purified naive P14 CD8^{+} T cells were resuspended in serum free DMEM and transferred by retro-orbital injection. For scRNA-seq experiments during acute and chronic LCMV infection, 2x10^{4} naive P14 CD8^{+} T cells were transferred per recipient mouse. For *Il2ra*$^{-/-}$ single cell experiment, 2.4x10^{5} naive P14 CD8^{+} T cells were transferred per recipient mouse (Thy1.1^{+} *Il2ra*$^{-/-}$ to Thy1.1^{+} Thy1.2^{+} ratio = 2:1). For retroviral transduction, purified naive CD8^{+} T cells were activated with anti-CD3 and anti-CD28, retrovirus was packaged and cells were transduced as described [25] with the following modifications. Naive CD8 T cells were activated for 16-18 hours, transduced for 4 hours with retroviral supernatants, and immediately transferred to naive 6-8 week old C57BL/6J hosts. 50,000 cells were transferred to each host and were infected with 2x10^{5} PFU of LCMV$_{Arm}$, or 500,000 cells were transferred and hosts were administered IP injection of 1.5x10^{5} PFU of LCMV$_{Cl13}$, or 1x10^{4} CFU of LM$_{GP33}$. LCMV$_{Arm}$, LCMV$_{Cl13}$ and LM$_{GP33}$ stocks were produced as described [25]. Infections were administered ~1 hour after adoptive transfer of transduced T cells, or the following day(s) after naive cell transfer. Virus stocks were stored at -80°C and thawed immediately before dilution. IP injection of 2x10^{5}

PFU of $LCMV_{Arm}$ per mouse was used to initiate acute infection, and retroorbital injection of $2x10^6$ PFU $LCMV_{Cl13}$ per mouse was used to initiate chronic infection.

Flow cytometry analysis and Sorting for single cell sequencing

Single cell suspensions were prepared by disrupting spleen sections by pressing through 70µm cell strainer in DMEM with 10% FBS. The splenocyte suspensions or heparnized peripheral blood collections were pelleted and red blood cells were lysed using ACK buffer. Cells were resuspended in FACS buffer, stained for surface proteins and then fixed in 2-4% PFA and permeabilzed for intracellular staining. Anti-mouse CD4 (RM4-5), CD8 (53-6.7), CD44 (IM7), CD62L (MEL-14), CD25 (3C7), CD127 (A7R34), KLRG1 (2F1/KLRG1), CD27 (LG.3A10), TCF-1 (S33-966), CX3CR1 (SA011F11) and were purchased from Biolegend or BD Biosciences. Intracellular staining for TCF-1 was performed using the Foxp3 transcription factor staining kit (eBioscience). For analysis of cells on days 3 or 4 of infection, spleens were cut into 1-2mm pieces and digested with collagenase IV (100 U/mL, Worthington) and DNase I (10 µg/mL, Sigma) in complete T cell media for 10 min at 37°C on a nutator, then disrupted over a 70µm cell strainer. For FACS isolation, $CD8^+$ T cells were initially enriched from total splenocyte preparations by negative selection with anti-CD4, anti-CD19, anti-B220 and anti-TER119 and magnetic streptavidin beads (Stemcell Technologies). Enriched cells were pre-stained with GP33-AF488 tetramer (NIH tetramer facility) to label endogenous LCMV-specific $CD8^+$ T cells, followed by staining with CD8 (BV421), Thy1.1 (PE) and Thy1.2 (APC) to label donor P14 $CD8^+$ T cells and the cells were sorted using a BD FACS Aria™ Fusion.

In vivo Pooled RNAi Screen

The RNAi screen was performed and analyzed as shown in fig S4A-B and as previously described [14] with the following modifications and details. Naive P14 CD8 T cells were activated for 18 hours using plate bound anti-CD3 and soluble anti-CD28 and transduced for 4 hours in 96 well plates before cells from all wells were pooled. Immediately after pooling (~24hrs post activation) aliquots of 500,000 cells were transferred into multiple naive host mice that were rested for ~1 hour after receiving cells before inoculation with $1.5x10^5$ PFU $LCMV_{Cl13}$ per mouse, which induces an acute infection in this setting[14]. Two entire biological replicates of the screen were performed and used for compuationatl analysis, and each replicate was screened in three pools. Each pool of shRNAmirs targeting CRFs was generated from ~500 shRNAmirs shRNAmirs which also included a common set of control shRNAmirs. Each pool was analyzed

in 10 recipient mice to maintain 50-100-fold theoretical representation of each shRNAmir after engraftment. For input samples, ~8x10^5 transduced cells were FACS-purified 48 hours after transduction. Eight days following infection, 3-8x10^5 cells from each KLRG1/CD127 gate were recovered by FACS from the spleens of infected mice. Genomic DNA was purified from each sample and used as PCR template to generate libraries for high-throughput sequencing as described [14]. Sequencing reads are mapped to library fasta file containing shRNA sequence information with custom blast pipeline. Raw read counts for each shRNA are normalized to total counts, and quantiles of each shRNA were calculated with negative binomial distribution. To calculate the effect size of shRNA in different cell population, the difference between quantile in different cell population was calculated for individual shNRAs. To sum up the effect of each gene, the quantile shift for all shRNAs towards each gene were converted to Z score, averaged and adjusted by p-value (to account for the consistency of effect) to generate the adjsusted Z-Scores for ranking (fig S4B).

Nascent RNA-seq analysis

For nascent RNA-seq of *in vitro* activated CD8 T cells, chromatin-associated RNA was prepared and total RNA-seq libraries were prepared and sequenced as described [25]. Pair end fastq reads were trimmed with “trim_galore --paired --length 24 --stringency 3”. Trimmed fastq reads were aligned to GRCm38 genome with bowtie2 with parameters “-p 16 -X 2000 --fr” [57]. Forward and reverse strand reads were separated with samtools [58]. Reads per transcript were counted with subread featureCounts [59]. Counts from forward and reverse strands were merged with custom python script. Differential analysis was performed with DEseq2 [60].

Single cell RNA-seq library generation, sequencing and analysis

To prepare barcoded scRNA-seq libriaries from multiple libraries, anti-mouse mouse MHC H-2 hashtag antibodies (Biolegend TotalSeq^{TM}) were used to label separate FACS-purified subsets which were washed, counted and pooled to final concentration of ~1,600 cells/ul. A total of ~50,000 cells were loaded into one lane of the single cell A chip kit (P/N 1000009). Single-cell gel beads in emulsion (GEM) were generated using the 10X Chromium single cell controller (10X Genomics, Pleasanton, CA). Single-cell GEM’s and sequencing libraries were generated using the Single-cell 3’ library and gel bead kit V2 (P/N 120267) according to manufacturer recommendations. The final library size distributions and amounts

were assessed using bioanalyzer analysis and further quantified with the NEBNext library quantification kit (P/N E7630). The cDNA and HTO libraries were pooled 10:1 and sequenced to a depth of 50,000 reads per cell for the cDNA and 2,000 reads per cell for the HTO library. Libraries were sequenced on the Illumina NovaSeq with the following 10X read format; Read 1 25bp, index i7 8bp, and Read 2 98bp. Around 500 - 1500 million reads were generated per experiment, yielding 60-90% sequencing saturation and around 1500-3000 median genes per cell after alignment. For hashtag library sequences, 60-75% antibody reads were mapped to cells (usable), yielding around 3000-6000 usable antibody reads per cell.

Cellranger 3.0 was used for fastq generation (cellranger mkfastq) and counting (cellranger count). Standard quality filtering was performed with Scanpy to remove genes expressed in less than 3 cells, doublets or low read cells (with gene / count per cell cutoff), and cells with high mitochondria count percentage [61]. Demultiplexing of Biolegend hashtags was performed with a custom python script to exclude doublets and dropouts. After quality filtering, reads were normalized to 10,000 per cell and logarithmized. Highly variable genes were identified with scanpy.pp.highly_variable_genes and selected. Count matrices were then regressed and scaled with scanpy.pp.regress_out and scanpy.pp.scale. UMAP projection was generated after PCA (scanpy.tl.pca) and nearest neighbor analysis (scanpy.pp.neighbors). Louvain cluster extraction (scanpy.tl.louvain) was performed on UMAP projection and the extracted clusters were analyzed with PAGA (scanpy.pl.paga) for cluster correlations [18] [13]. For velocity-based analysis, Velocyto was used to generate spliced / unspliced count matrix [12]. The resulting matrix was processed in scVelo for velocity analysis [11]. For correlation of independent scRNA-seq experiments, Harmonypy was used for batch effect removal on normalized and scaled count matrices of experiments [62]. Dimensionality reduction was performed on batch effect removed matrices with PCA, followed by UMAP projection.

ChIP-seq analysis

Raw fastq files of ChIP-seq experiments were downloaded from GEO with SRA-Tools fastq-dump. Trimming of fastq files were performed with trim_galore. Trimpped reads were aligned to GRCm38 genome with bowtie2 [57]. Aligned sam file were sorted and filtered for PCR duplicates with samtools. Blacklisted regions were filtered with bedtools [63,64]. Peaks were called with MACS2 and annotated with R package ChIPSeeker [65,66]. All analytical codes for ChIP-seq are published on Github: https://github.com/TCellResearchTeam/T_Cell_ChIP. Visualization of ChIP data is accessible on UCSC Track Hub: T_cell_ATAC_ChIP_Pipkin.

GSEA

GSEA was performed with R package clusterProfiler [20]. GSEA signatures were downloaded or generated from published datasets available from GEO database. All signatures and description are available in https://github.com/TCellResearchTeam/T_cell_signature_Reference.

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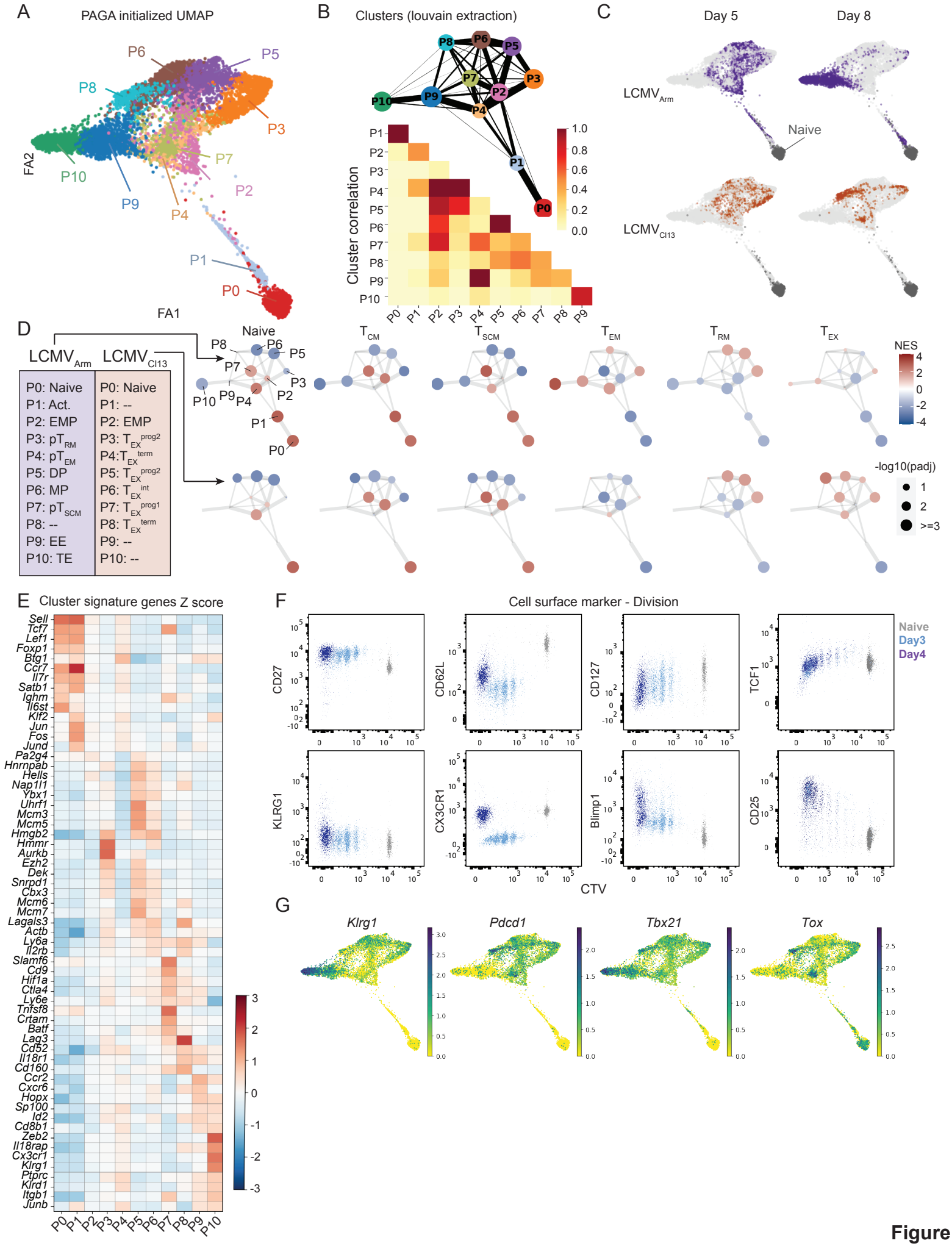

A
PAGA initialized UMAP
FA1
FA2
P0
P1
P2
P3
P4
P5
P6
P7
P8
P9
P10
B
Clusters (louvain extraction)
Cluster correlation
C
Day 5
Day 8
LCMV$_{Arm}$
LCMV$_{Cl13}$
Naive
D
LCMV$_{Arm}$
LCMV$_{Cl13}$
P0: Naive
P1: Act.
P2: EMP
P3: pT$_{RM}$
P4: pT$_{EM}$
P5: DP
P6: MP
P7: pT$_{SCM}$
P8: --
P9: EE
P10: TE
P0: Naive
P1: --
P2: EMP
P3: T$_{EX}^{prog2}$
P4: T$_{EX}^{term}$
P5: T$_{EX}^{prog2}$
P6: T$_{EX}^{int}$
P7: T$_{EX}^{prog1}$
P8: T$_{EX}^{term}$
P9: --
P10: --
T$_{CM}$
T$_{SCM}$
T$_{EM}$
T$_{RM}$
T$_{EX}$
NES
-log10(padj)
E
Cluster signature genes Z score
Sell
Tcf7
Lef1
Foxp1
Btg1
Ccr7
Il7r
Satb1
Ighm
Il6st
Klf2
Jun
Fos
Jund
Pa2g4
Hnrnpab
Hells
Nap1l1
Ybx1
Uhrf1
Mcm3
Mcm5
Hmgb2
Hmmr
Aurkb
Ezh2
Dek
Snrpd1
Cbx3
Mcm6
Mcm7
Lagals3
Actb
Ly6a
Il2rb
Slamf6
Cd9
Hif1a
Ctla4
Ly6e
Tnfsf8
Crtam
Batf
Lag3
Cd52
Il18r1
Cd160
Ccr2
Cxcr6
Hopx
Sp100
Id2
Cd8b1
Zeb2
Il18rap
Cx3cr1
Klrg1
Ptprc
Klrd1
Itgb1
Junb
F
Cell surface marker - Division
CD27
CD62L
CD127
TCF1
KLRG1
CX3CR1
Blimp1
CD25
CTV
Naive
Day3
Day4
G
Klrg1
Pdcd1
Tbx21
Tox

Figure 1

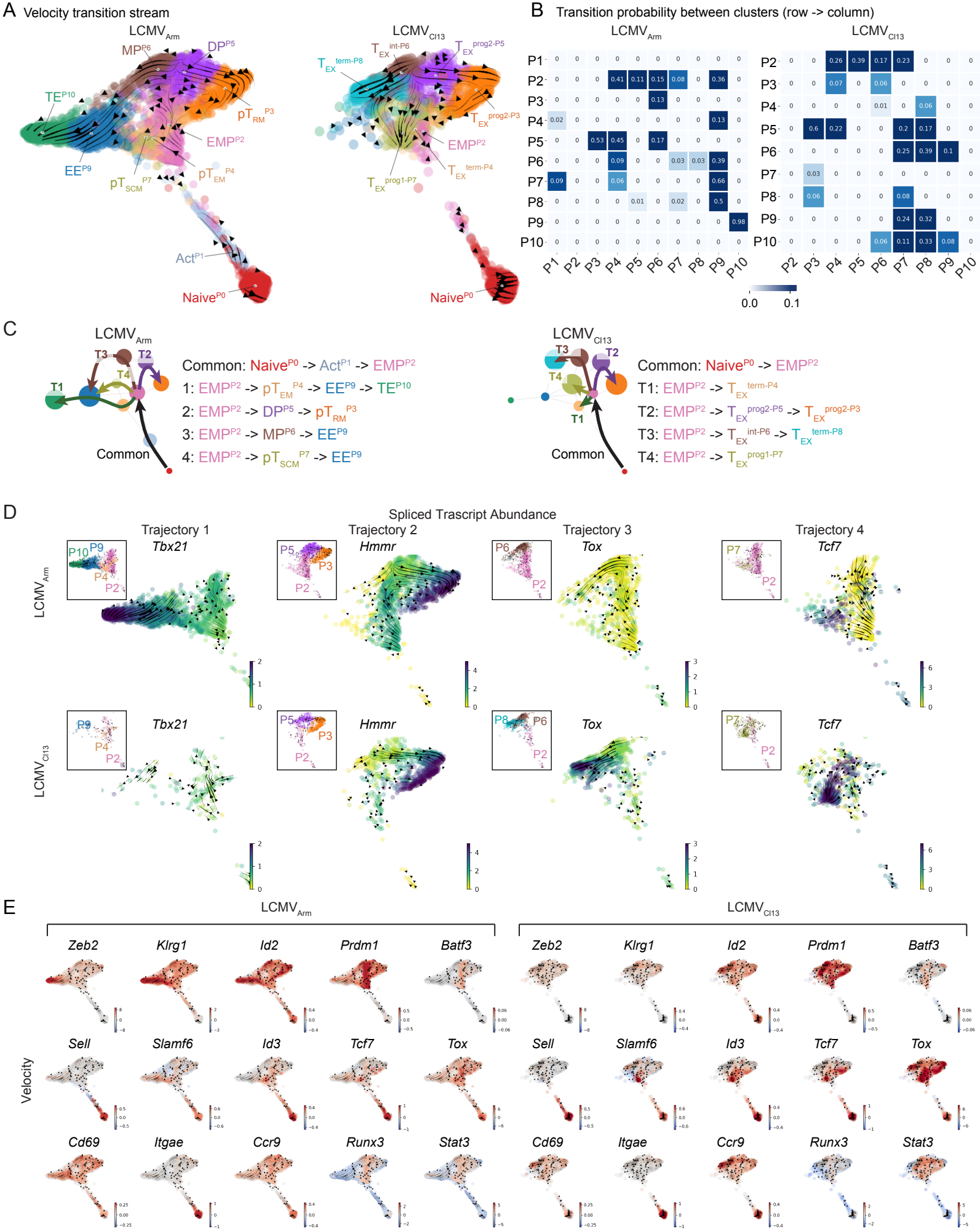

A Velocity transition stream
LCMV Arm
LCMV Cl13
B Transition probability between clusters (row -> column)
C
Common: Naive P0 -> Act P1 -> EMP P2
1: EMP P2 -> pT EM P4 -> EE P9 -> TE P10
2: EMP P2 -> DP P5 -> pT RM P3
3: EMP P2 -> MP P6 -> EE P9
4: EMP P2 -> pT SCM P7 -> EE P9
Common: Naive P0 -> EMP P2
T1: EMP P2 -> T EX term-P4
T2: EMP P2 -> T EX prog2-P5 -> T EX prog2-P3
T3: EMP P2 -> T EX int-P6 -> T EX term-P8
T4: EMP P2 -> T EX prog1-P7
D
Spliced Trascript Abundance
Trajectory 1
Trajectory 2
Trajectory 3
Trajectory 4
Tbx21
Hmmr
Tox
Tcf7
E
Velocity
Zeb2
Klrg1
Id2
Prdm1
Batf3
Sell
Slamf6
Id3
Tcf7
Tox
Cd69
Itgae
Ccr9
Runx3
Stat3

Figure 2

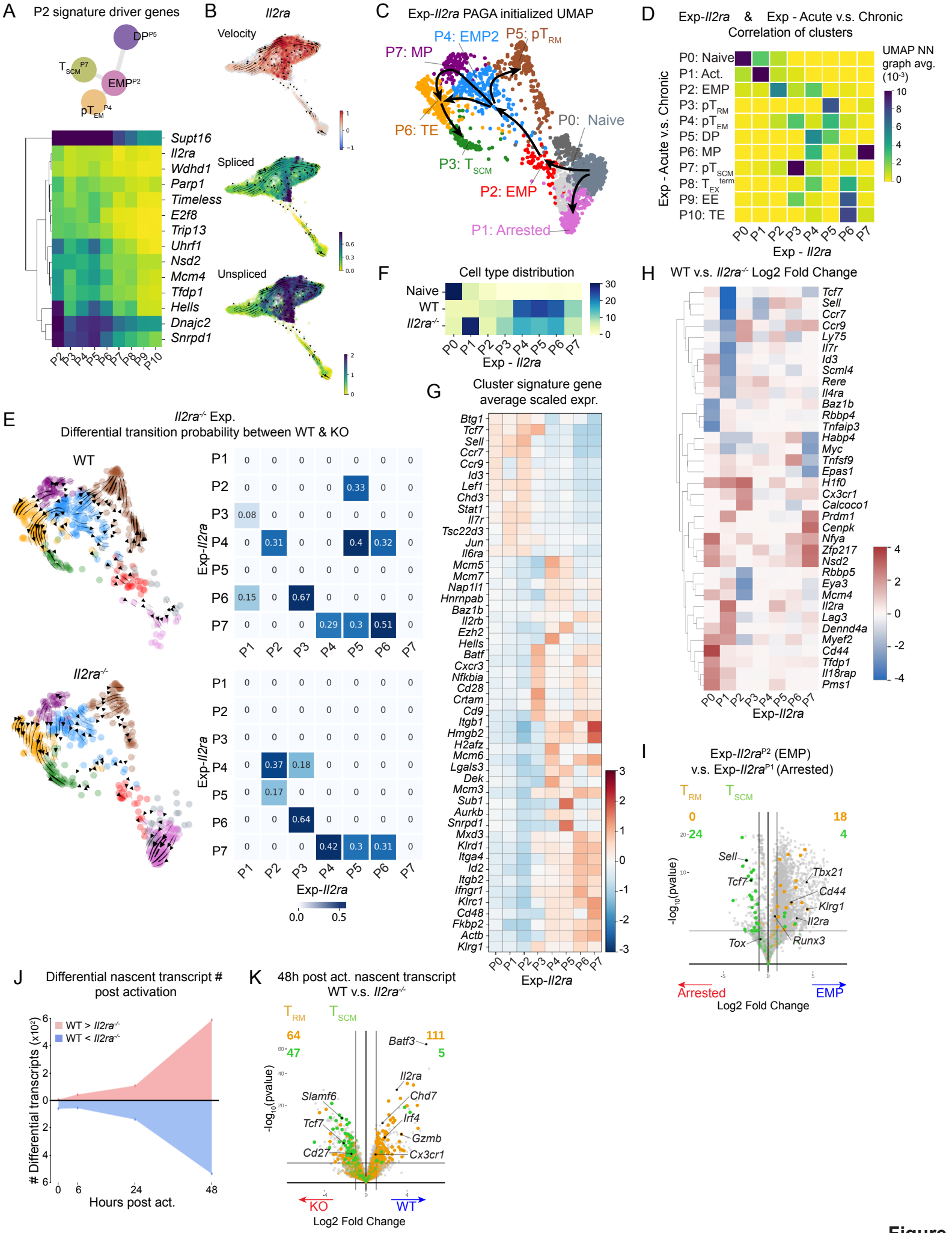

A
P2 signature driver genes
DP P5
T SCM P7
EMP P2
pT EM P4
Supt16
Il2ra
Wdhd1
Parp1
Timeless
E2f8
Trip13
Uhrf1
Nsd2
Mcm4
Tfdp1
Hells
Dnajc2
Snrpd1
P2 P3 P4 P5 P6 P7 P8 P9 P10
B
Il2ra
Velocity
Spliced
Unspliced
C
Exp-Il2ra PAGA initialized UMAP
P4: EMP2
P5: pT RM
P7: MP
P6: TE
P0: Naive
P3: T SCM
P2: EMP
P1: Arrested
D
Exp-Il2ra & Exp - Acute v.s. Chronic
Correlation of clusters
UMAP NN graph avg. (10-3)
P0: Naive
P1: Act.
P2: EMP
P3: pT RM
P4: pT EM
P5: DP
P6: MP
P7: pT SCM
P8: T EX term
P9: EE
P10: TE
Exp - Acute v.s. Chronic
Exp - Il2ra
E
Il2ra-/- Exp.
Differential transition probability between WT & KO
WT
Il2ra-/-
Exp-Il2ra
F
Cell type distribution
Naive
WT
Il2ra-/-
Exp - Il2ra
G
Cluster signature gene
average scaled expr.
Exp-Il2ra
H
WT v.s. Il2ra-/- Log2 Fold Change
Exp-Il2ra
I
Exp-Il2ra P2 (EMP)
v.s. Exp-Il2ra P1 (Arrested)
T RM
T SCM
Sell
Tcf7
Tbx21
Cd44
Klrg1
Il2ra
Tox
Runx3
-log10(pvalue)
Arrested
EMP
Log2 Fold Change
J
Differential nascent transcript #
post activation
WT > Il2ra-/-
WT < Il2ra-/-
Differential transcripts (x10²)
Hours post act.
K
48h post act. nascent transcript
WT v.s. Il2ra-/-
T RM
T SCM
Batf3
Il2ra
Chd7
Slamf6
Tcf7
Irf4
Gzmb
Cd27
Cx3cr1
-log10(pvalue)
KO
WT
Log2 Fold Change

Figure 3

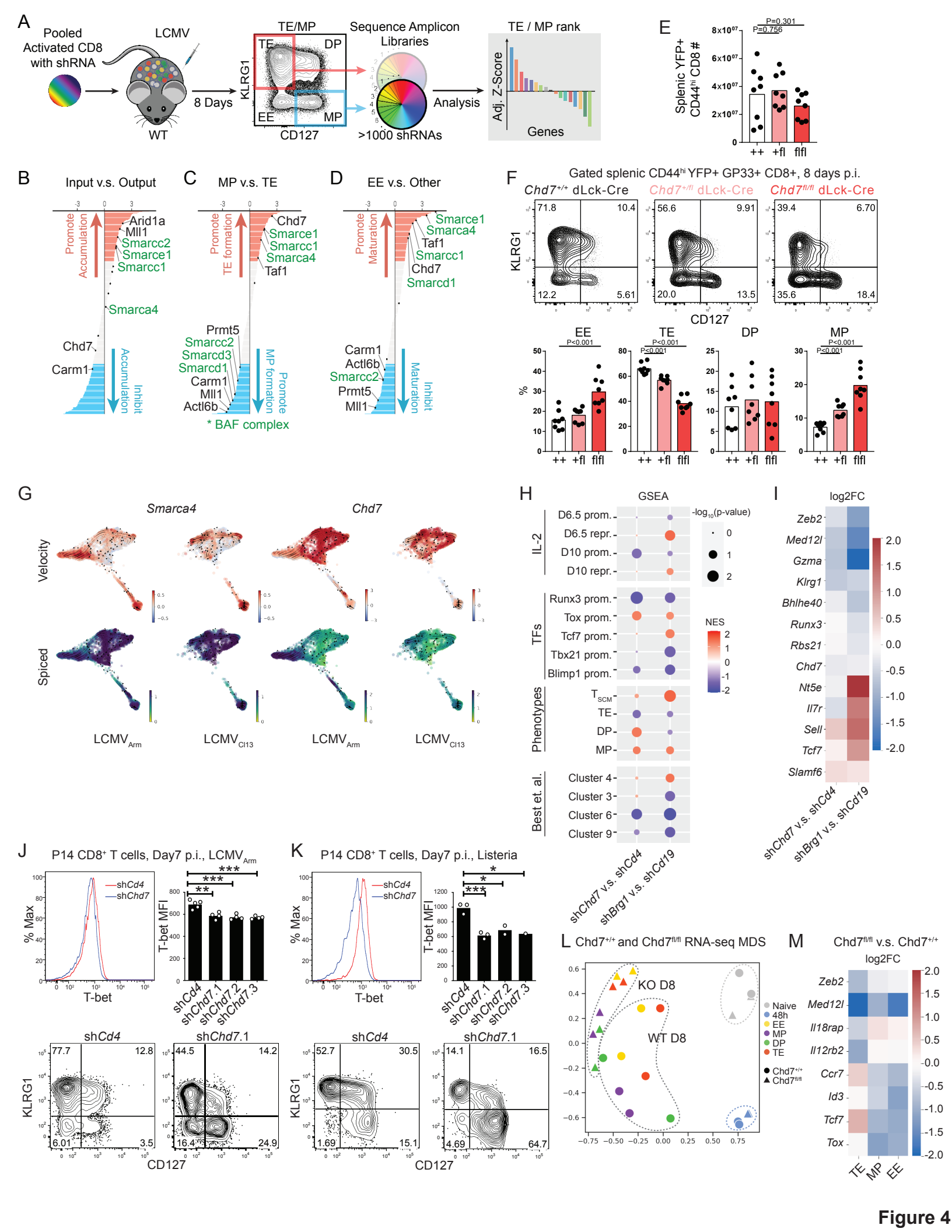

A
Pooled Activated CD8 with shRNA
LCMV
WT
8 Days
TE/MP
TE
DP
EE
MP
KLRG1
CD127
Sequence Amplicon Libraries
>1000 shRNAs
Analysis
TE / MP rank
Adj. Z-Score
Genes
E
Splenic YFP+ CD44hi CD8 #
P=0.301
P=0.756
++
+fl
flfl
B
Input v.s. Output
Promote Accumulation
Arid1a
Mll1
Smarcc2
Smarce1
Smarcc1
Smarca4
Chd7
Carm1
Accumulation Inhibit
C
MP v.s. TE
Promote TE formation
Chd7
Smarce1
Smarcc1
Smarca4
Taf1
Prmt5
Smarcc2
Smarcd3
Smarcd1
Carm1
Mll1
Actl6b
MP formation Promote
* BAF complex
D
EE v.s. Other
Promote Maturation
Smarce1
Smarca4
Taf1
Smarcc1
Chd7
Smarcd1
Carm1
Actl6b
Smarcc2
Prmt5
Mll1
Maturation Inhibit
F
Gated splenic CD44hi YFP+ GP33+ CD8+, 8 days p.i.
Chd7+/+ dLck-Cre
Chd7+/fl dLck-Cre
Chd7fl/fl dLck-Cre
71.8
10.4
12.2
5.61
56.6
9.91
20.0
13.5
39.4
6.70
35.6
18.4
EE
TE
DP
MP
P<0.001
%
G
Smarca4
Chd7
Velocity
Spiced
LCMVArm
LCMVCl13
H
GSEA
-log10(p-value)
IL-2
D6.5 prom.
D6.5 repr.
D10 prom.
D10 repr.
TFs
Runx3 prom.
Tox prom.
Tcf7 prom.
Tbx21 prom.
Blimp1 prom.
Phenotypes
TSCM
TE
DP
MP
Best et. al.
Cluster 4
Cluster 3
Cluster 6
Cluster 9
NES
shChd7 v.s. shCd4
shBrg1 v.s. shCd19
I
log2FC
Zeb2
Med12l
Gzma
Klrg1
Bhlhe40
Runx3
Rbs21
Chd7
Nt5e
Il7r
Sell
Tcf7
Slamf6
shChd7 v.s. shCd4
shBrg1 v.s. shCd19
J
P14 CD8+ T cells, Day7 p.i., LCMVArm
shCd4
shChd7
% Max
T-bet
T-bet MFI
shCd4
shChd7.1
shChd7.2
shChd7.3
77.7
12.8
6.01
3.5
44.5
14.2
16.4
24.9
KLRG1
CD127
K
P14 CD8+ T cells, Day7 p.i., Listeria
52.7
30.5
1.69
15.1
14.1
16.5
4.69
64.7
L
Chd7+/+ and Chd7fl/fl RNA-seq MDS
KO D8
WT D8
Naive
48h
EE
MP
DP
TE
Chd7+/+
Chd7fl/fl
M
Chd7fl/fl v.s. Chd7+/+
log2FC
Zeb2
Med12l
Il18rap
Il12rb2
Ccr7
Id3
Tcf7
Tox
TE
MP
EE

Figure 4

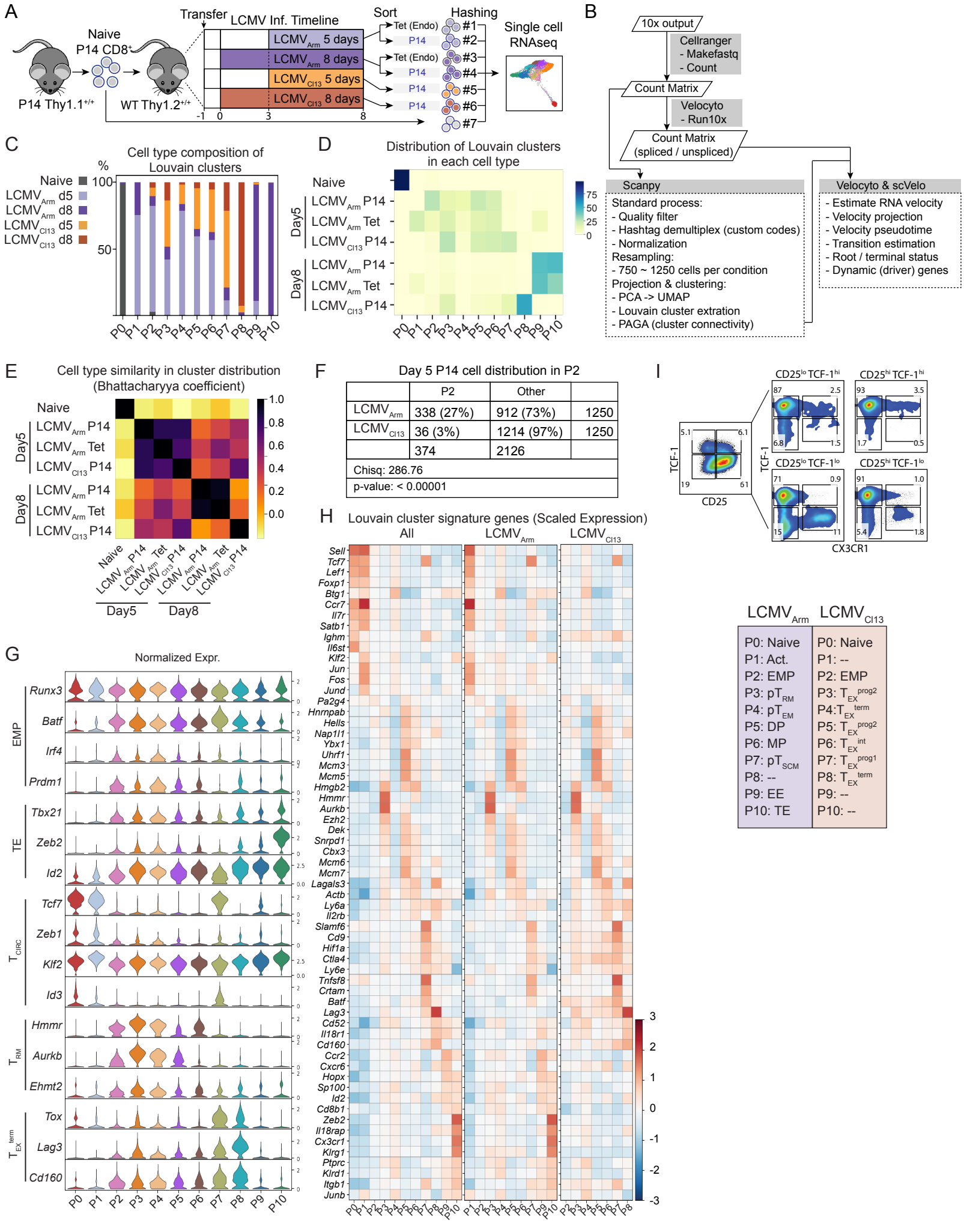

A
Naive P14 CD8+
P14 Thy1.1+/+
WT Thy1.2+/+
Transfer
LCMV Inf. Timeline
LCMVArm 5 days
LCMVArm 8 days
LCMVCl13 5 days
LCMVCl13 8 days
Sort
Tet (Endo)
P14
Hashing
#1
#2
#3
#4
#5
#6
#7
Single cell RNAseq
B
10x output
Cellranger
- Makefastq
- Count
Count Matrix
Velocyto
- Run10x
Count Matrix (spliced / unspliced)
Scanpy
Standard process:
- Quality filter
- Hashtag demultiplex (custom codes)
- Normalization
Resampling:
- 750 ~ 1250 cells per condition
Projection & clustering:
- PCA -> UMAP
- Louvain cluster extration
- PAGA (cluster connectivity)
Velocyto & scVelo
- Estimate RNA velocity
- Velocity projection
- Velocity pseudotime
- Transition estimation
- Root / terminal status
- Dynamic (driver) genes
C
Cell type composition of Louvain clusters
Naive
LCMVArm d5
LCMVArm d8
LCMVCl13 d5
LCMVCl13 d8
D
Distribution of Louvain clusters in each cell type
Day5
Day8
E
Cell type similarity in cluster distribution (Bhattacharyya coefficient)
F
Day 5 P14 cell distribution in P2
P2
Other
LCMVArm 338 (27%) 912 (73%) 1250
LCMVCl13 36 (3%) 1214 (97%) 1250
374 2126
Chisq: 286.76
p-value: < 0.00001
G
Normalized Expr.
EMP
TE
TCIRC
TRM
TEXterm
H
Louvain cluster signature genes (Scaled Expression)
All
LCMVArm
LCMVCl13
I
CD25lo TCF-1hi
CD25hi TCF-1hi
CD25lo TCF-1lo
CD25hi TCF-1lo
TCF-1
CD25
CX3CR1
LCMVArm
LCMVCl13
P0: Naive
P1: Act.
P2: EMP
P3: pTRM
P4: pTEM
P5: DP
P6: MP
P7: pTSCM
P8: --
P9: EE
P10: TE
P0: Naive
P1: --
P2: EMP
P3: TEXprog2
P4: TEXterm
P5: TEXprog2
P6: TEXint
P7: TEXprog1
P8: TEXterm
P9: --
P10: --

Figure S1

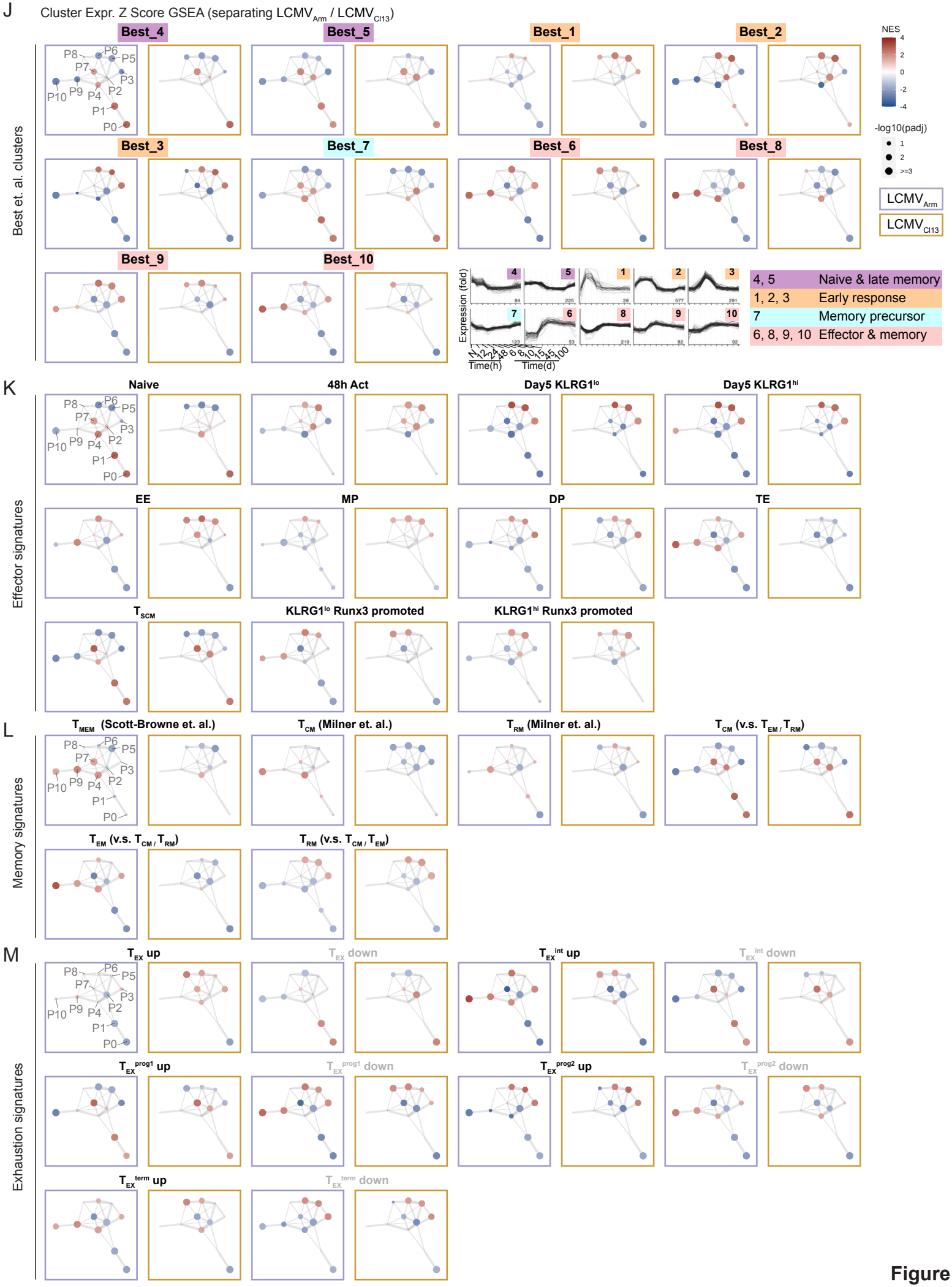

J
Cluster Expr. Z Score GSEA (separating LCMV$_{Arm}$ / LCMV$_{Cl13}$)
Best et. al. clusters
Best_4
Best_5
Best_1
Best_2
Best_3
Best_7
Best_6
Best_8
Best_9
Best_10
P8 P6 P5 P7 P3 P10 P9 P4 P2 P1 P0
NES
4 2 0 -2 -4
-log10(padj)
1 2 >=3
LCMV$_{Arm}$
LCMV$_{Cl13}$
Expression (fold)
Time(h)
Time(d)
4, 5 Naive & late memory
1, 2, 3 Early response
7 Memory precursor
6, 8, 9, 10 Effector & memory
K
Effector signatures
Naive
48h Act
Day5 KLRG1^{lo}
Day5 KLRG1^{hi}
EE
MP
DP
TE
T$_{SCM}$
KLRG1^{lo} Runx3 promoted
KLRG1^{hi} Runx3 promoted
L
Memory signatures
T$_{MEM}$ (Scott-Browne et. al.)
T$_{CM}$ (Milner et. al.)
T$_{RM}$ (Milner et. al.)
T$_{CM}$ (v.s. T$_{EM}$ / T$_{RM}$)
T$_{EM}$ (v.s. T$_{CM}$ / T$_{RM}$)
T$_{RM}$ (v.s. T$_{CM}$ / T$_{EM}$)
M
Exhaustion signatures
T$_{EX}$ up
T$_{EX}$ down
T$_{EX}^{int}$ up
T$_{EX}^{int}$ down
T$_{EX}^{prog1}$ up
T$_{EX}^{prog1}$ down
T$_{EX}^{prog2}$ up
T$_{EX}^{prog2}$ down
T$_{EX}^{term}$ up
T$_{EX}^{term}$ down

Figure S1

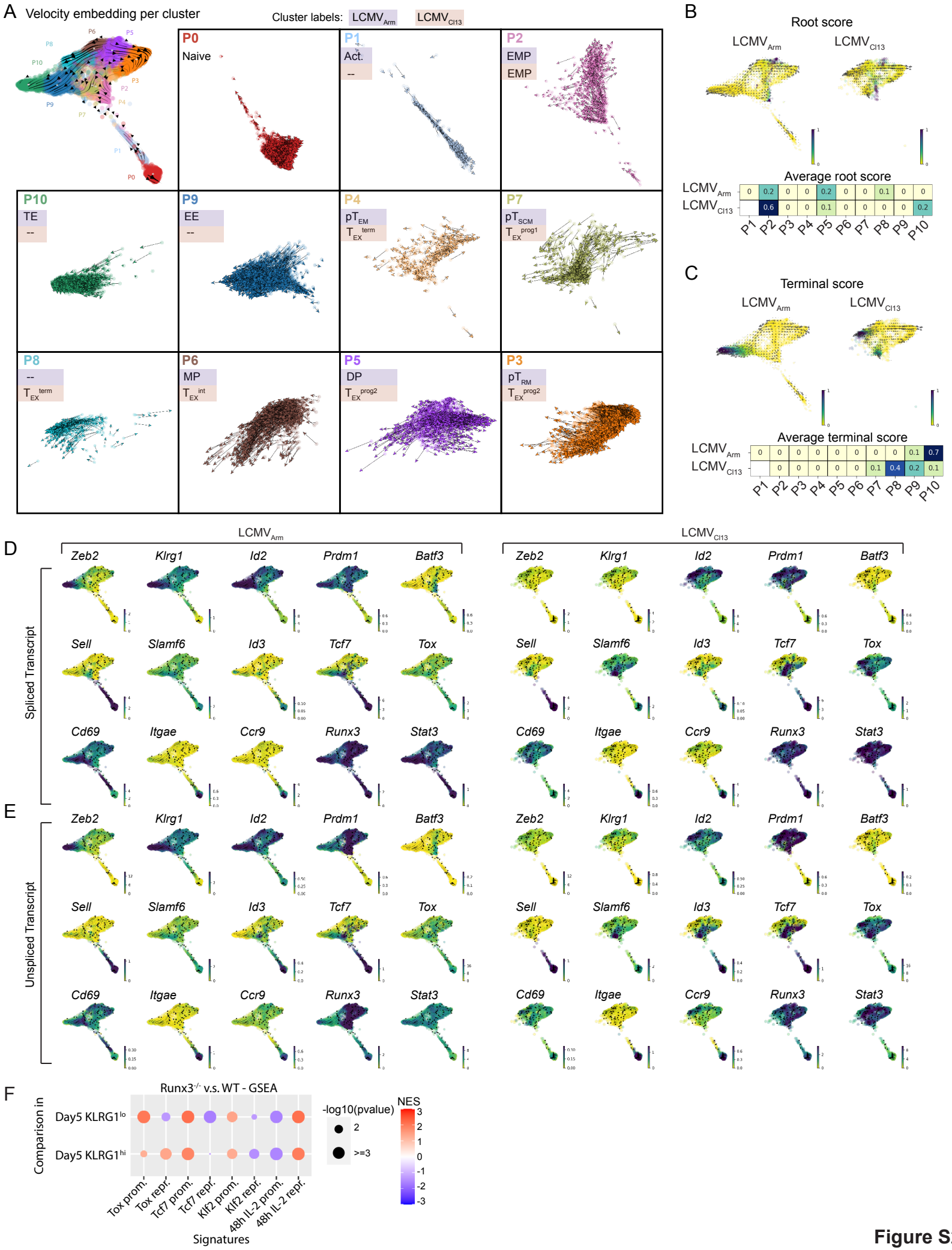

A
Velocity embedding per cluster
Cluster labels: LCMV Arm LCMV Cl13
P0 Naive
P1 Act. --
P2 EMP EMP
P10 TE --
P9 EE --
P4 pT EM T EX term
P7 pT SCM T EX prog1
P8 -- T EX term
P6 MP T EX int
P5 DP T EX prog2
P3 pT RM T EX prog2
B
Root score
LCMV Arm
LCMV Cl13
Average root score
C
Terminal score
Average terminal score
D
E
Spliced Transcript
Unspliced Transcript
Zeb2 Klrg1 Id2 Prdm1 Batf3
Sell Slamf6 Id3 Tcf7 Tox
Cd69 Itgae Ccr9 Runx3 Stat3
F
Runx3-/- v.s. WT - GSEA
Comparison in
Day5 KLRG1lo
Day5 KLRG1hi
Tox prom. Tox repr. Tcf7 prom. Tcf7 repr. Klf2 prom. Klf2 repr. 48h IL-2 prom. 48h IL-2 repr.
Signatures
-log10(pvalue)
NES

Figure S2

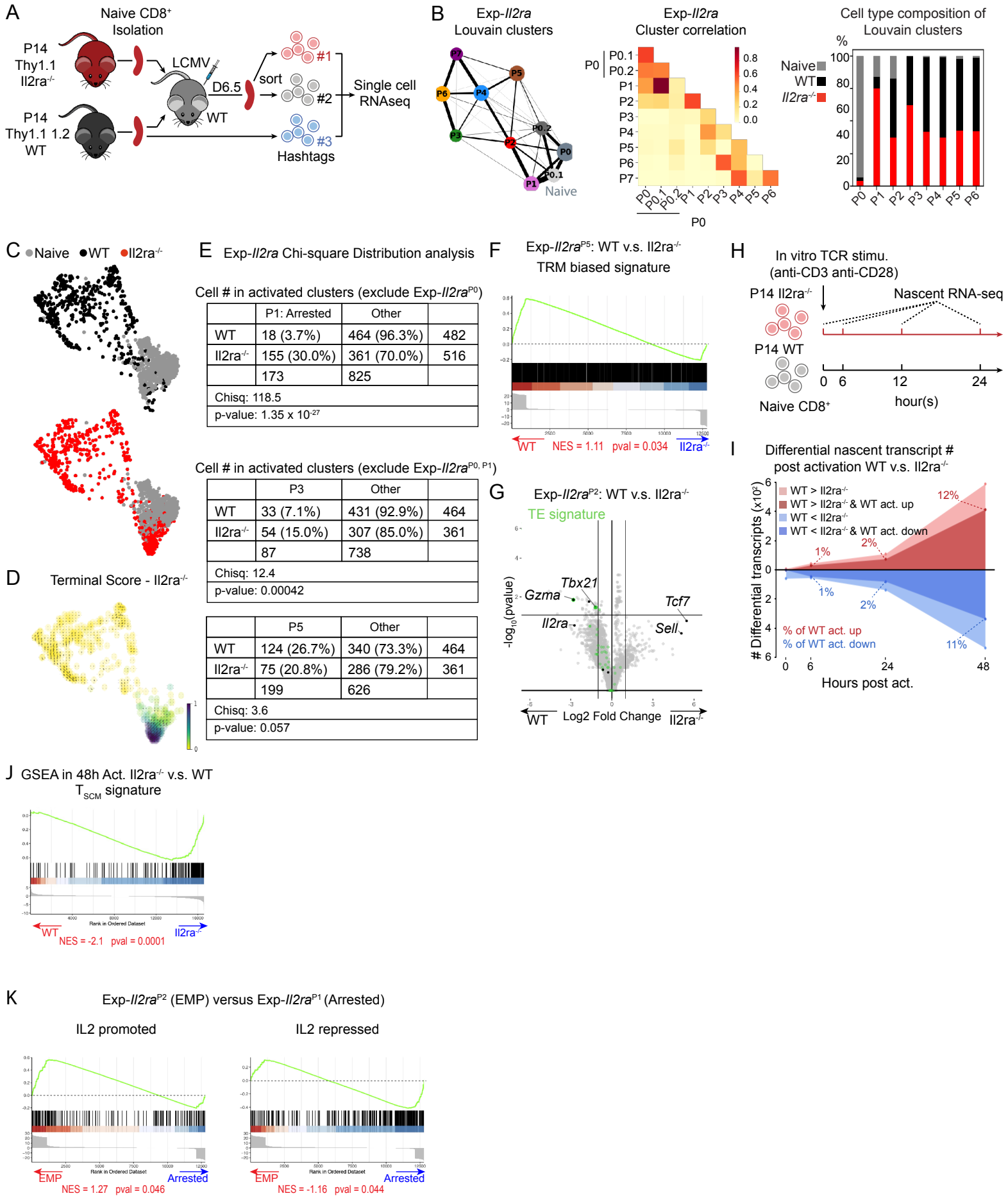

A
Naive CD8+ Isolation
P14 Thy1.1 Il2ra-/-
P14 Thy1.1 1.2 WT
LCMV
D6.5
WT
sort
#1
#2
#3
Hashtags
Single cell RNAseq
B
Exp-Il2ra Louvain clusters
Naive
Exp-Il2ra Cluster correlation
Cell type composition of Louvain clusters
Naive
WT
Il2ra-/-
C
Naive
WT
Il2ra-/-
D
Terminal Score - Il2ra-/-
E
Exp-Il2ra Chi-square Distribution analysis
Cell # in activated clusters (exclude Exp-Il2ra P0)
P1: Arrested
Other
WT 18 (3.7%) 464 (96.3%) 482
Il2ra-/- 155 (30.0%) 361 (70.0%) 516
173 825
Chisq: 118.5
p-value: 1.35 x 10-27
Cell # in activated clusters (exclude Exp-Il2ra P0, P1)
P3
Other
WT 33 (7.1%) 431 (92.9%) 464
Il2ra-/- 54 (15.0%) 307 (85.0%) 361
87 738
Chisq: 12.4
p-value: 0.00042
P5
Other
WT 124 (26.7%) 340 (73.3%) 464
Il2ra-/- 75 (20.8%) 286 (79.2%) 361
199 626
Chisq: 3.6
p-value: 0.057
F
Exp-Il2ra P5: WT v.s. Il2ra-/-
TRM biased signature
WT
NES = 1.11 pval = 0.034
Il2ra-/-
G
Exp-Il2ra P2: WT v.s. Il2ra-/-
TE signature
Tbx21
Gzma
Il2ra
Tcf7
Sell
-log10(pvalue)
WT
Log2 Fold Change
Il2ra-/-
H
In vitro TCR stimu. (anti-CD3 anti-CD28)
P14 Il2ra-/-
P14 WT
Naive CD8+
Nascent RNA-seq
hour(s)
I
Differential nascent transcript # post activation WT v.s. Il2ra-/-
WT > Il2ra-/-
WT > Il2ra-/- & WT act. up
WT < Il2ra-/-
WT < Il2ra-/- & WT act. down
% of WT act. up
% of WT act. down
# Differential transcripts (x10²)
Hours post act.
J
GSEA in 48h Act. Il2ra-/- v.s. WT
TSCM signature
Rank in Ordered Dataset
WT
NES = -2.1 pval = 0.0001
Il2ra-/-
K
Exp-Il2ra P2 (EMP) versus Exp-Il2ra P1 (Arrested)
IL2 promoted
IL2 repressed
EMP
Arrested
NES = 1.27 pval = 0.046
NES = -1.16 pval = 0.044

Figure S3

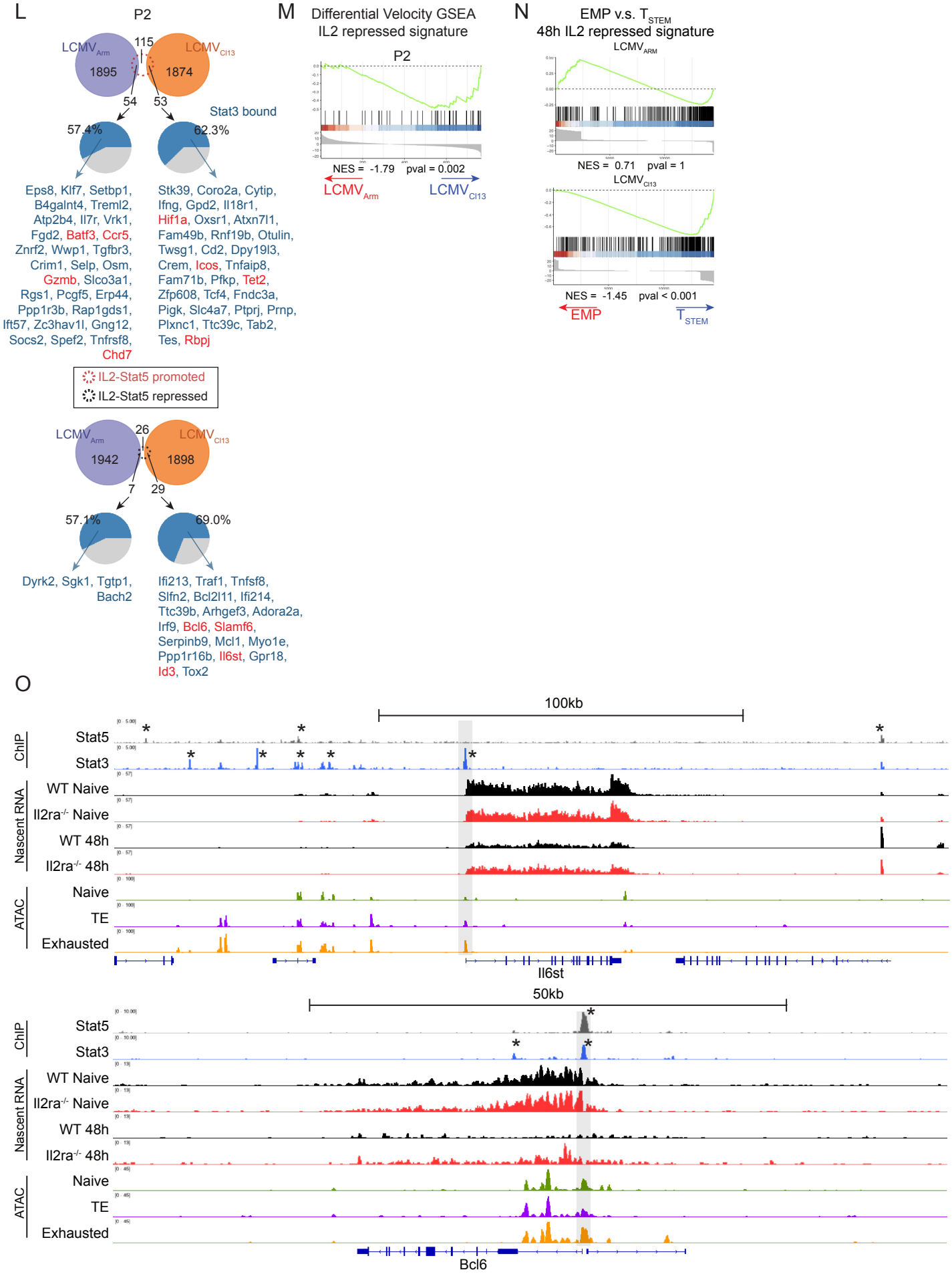

L
P2
LCMV Arm
LCMV Cl13
1895
1874
115
54
53
Stat3 bound
57.4%
62.3%
Eps8, Klf7, Setbp1, B4galnt4, Treml2, Atp2b4, Il7r, Vrk1, Fgd2, Batf3, Ccr5, Znrf2, Wwp1, Tgfbr3, Crim1, Selp, Osm, Gzmb, Slco3a1, Rgs1, Pcgf5, Erp44, Ppp1r3b, Rap1gds1, Ift57, Zc3hav1l, Gng12, Socs2, Spef2, Tnfrsf8, Chd7
Stk39, Coro2a, Cytip, Ifng, Gpd2, Il18r1, Hif1a, Oxsr1, Atxn7l1, Fam49b, Rnf19b, Otulin, Twsg1, Cd2, Dpy19l3, Crem, Icos, Tnfaip8, Fam71b, Pfkp, Tet2, Zfp608, Tcf4, Fndc3a, Pigk, Slc4a7, Ptprj, Prnp, Plxnc1, Ttc39c, Tab2, Tes, Rbpj
IL2-Stat5 promoted
IL2-Stat5 repressed
LCMV Arm
LCMV Cl13
1942
1898
26
7
29
57.1%
69.0%
Dyrk2, Sgk1, Tgtp1, Bach2
Ifi213, Traf1, Tnfsf8, Slfn2, Bcl2l11, Ifi214, Ttc39b, Arhgef3, Adora2a, Irf9, Bcl6, Slamf6, Serpinb9, Mcl1, Myo1e, Ppp1r16b, Il6st, Gpr18, Id3, Tox2
M
Differential Velocity GSEA
IL2 repressed signature
P2
NES = -1.79 pval = 0.002
LCMV Arm
LCMV Cl13
N
EMP v.s. T STEM
48h IL2 repressed signature
LCMV ARM
NES = 0.71 pval = 1
LCMV Cl13
NES = -1.45 pval < 0.001
EMP
T STEM
O
100kb
ChIP
Stat5
Stat3
Nascent RNA
WT Naive
Il2ra-/- Naive
WT 48h
Il2ra-/- 48h
ATAC
Naive
TE
Exhausted
Il6st
50kb
Bcl6

Figure S3

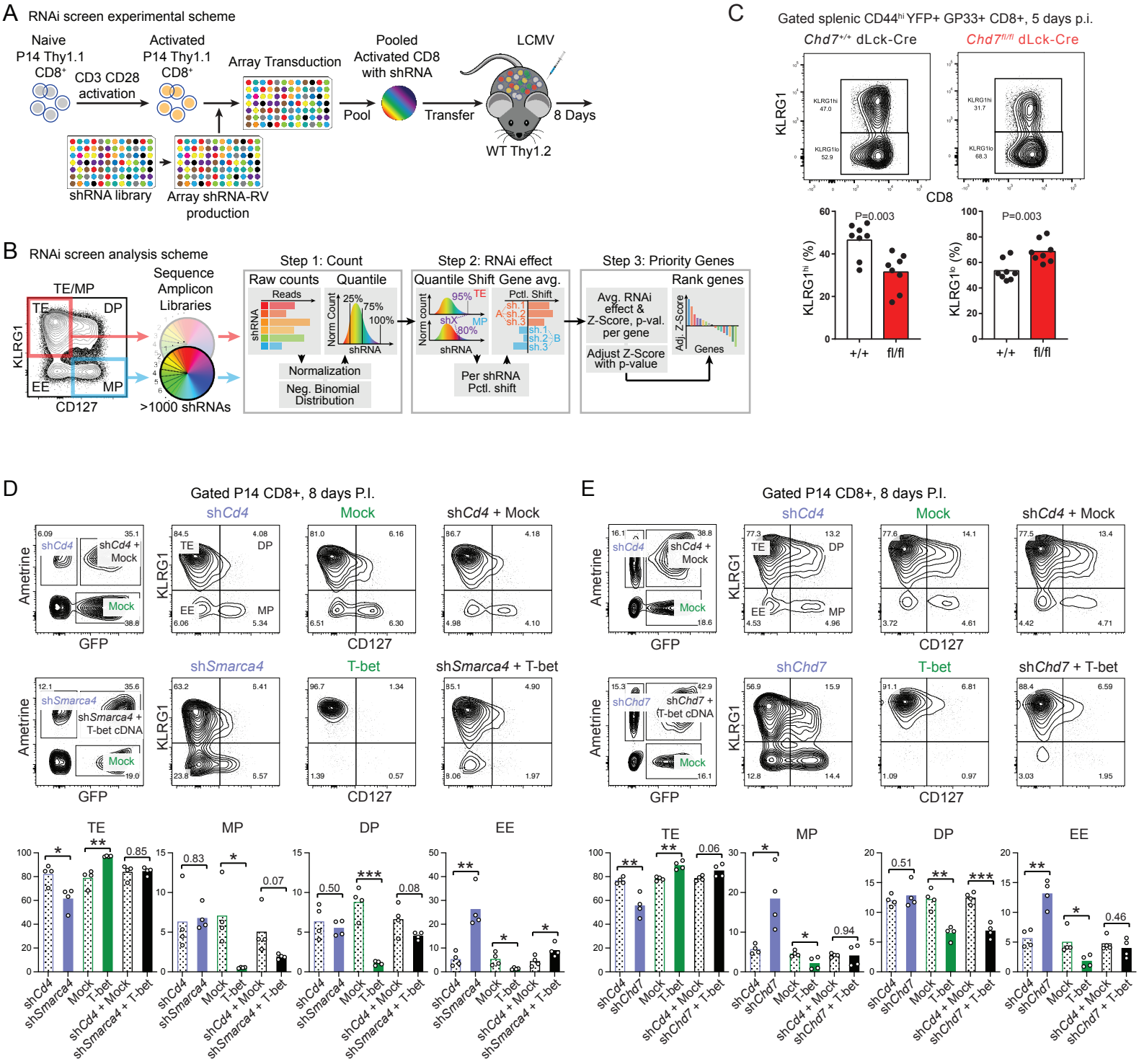

A
RNAi screen experimental scheme
Naive P14 Thy1.1 CD8+
CD3 CD28 activation
Activated P14 Thy1.1 CD8+
Array Transduction
Pooled Activated CD8 with shRNA
Pool
Transfer
LCMV
8 Days
WT Thy1.2
shRNA library
Array shRNA-RV production
B
RNAi screen analysis scheme
TE/MP
TE
DP
EE
MP
KLRG1
CD127
Sequence Amplicon Libraries
>1000 shRNAs
Step 1: Count
Raw counts
Quantile
Reads
shRNA
Norm. Count
25%
75%
100%
Normalization
Neg. Binomial Distribution
Step 2: RNAi effect
Quantile Shift Gene avg.
Norm count
95%
80%
Pctl. Shift
Per shRNA Pctl. shift
Step 3: Priority Genes
Avg. RNAi effect & Z-Score, p-val. per gene
Adjust Z-Score with p-value
Rank genes
Adj. Z-Score
Genes
C
Gated splenic CD44hi YFP+ GP33+ CD8+, 5 days p.i.
Chd7+/+ dLck-Cre
Chd7fl/fl dLck-Cre
KLRG1
CD8
KLRG1hi 47.0
KLRG1lo 52.9
KLRG1hi 31.7
KLRG1lo 68.3
P=0.003
KLRG1hi (%)
KLRG1lo (%)
+/+
fl/fl
D
Gated P14 CD8+, 8 days P.I.
shCd4
Mock
shCd4 + Mock
shSmarca4
T-bet
shSmarca4 + T-bet
Ametrine
GFP
CD127
shSmarca4 + T-bet cDNA
TE
MP
DP
EE
shCd4
shSmarca4
Mock
T-bet
shCd4 + Mock
shSmarca4 + T-bet
E
Gated P14 CD8+, 8 days P.I.
shCd4
Mock
shCd4 + Mock
shChd7
T-bet
shChd7 + T-bet
shCd4 + Mock
shChd7 + T-bet
shChd7 + T-bet cDNA

Figure S4

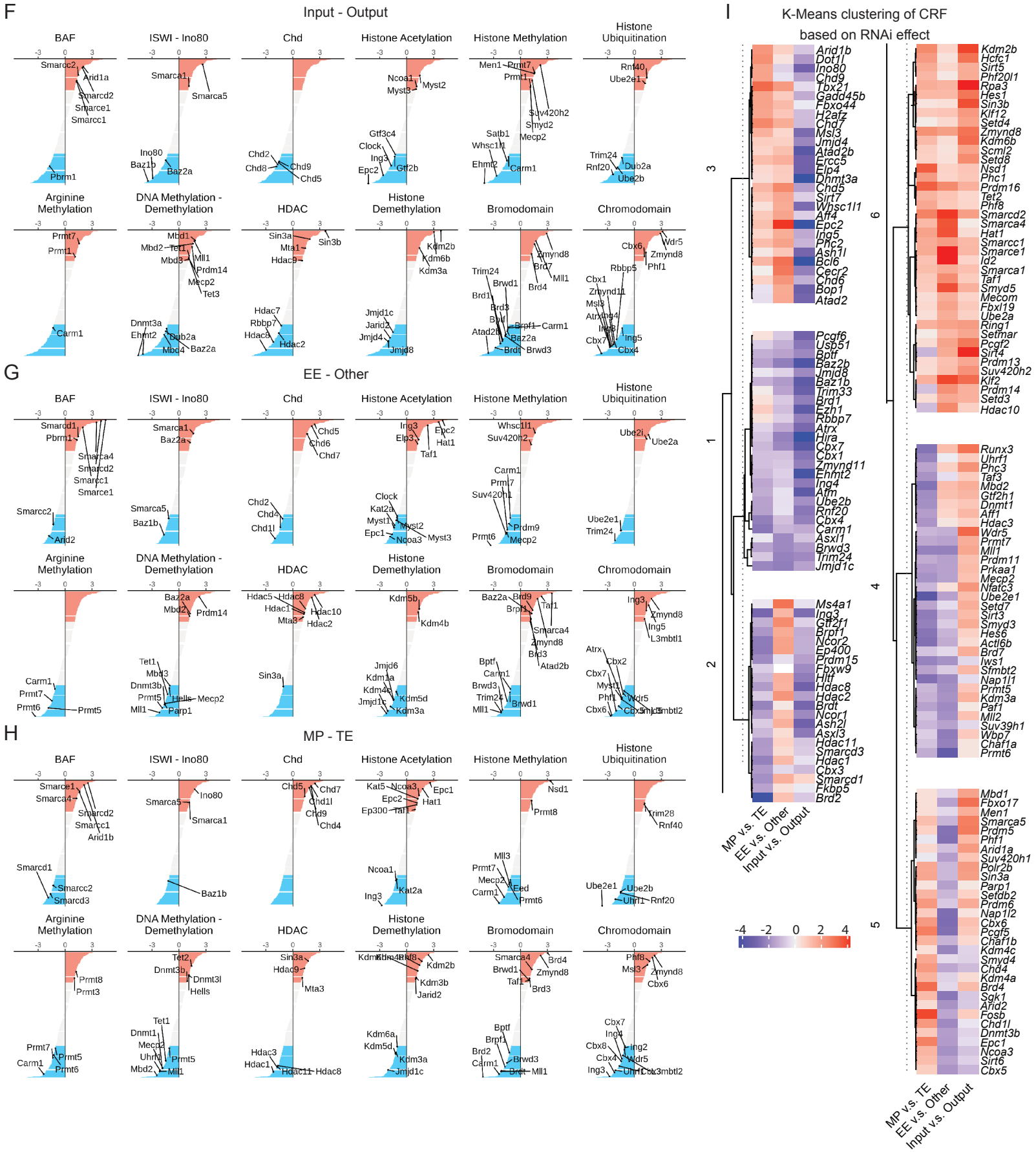
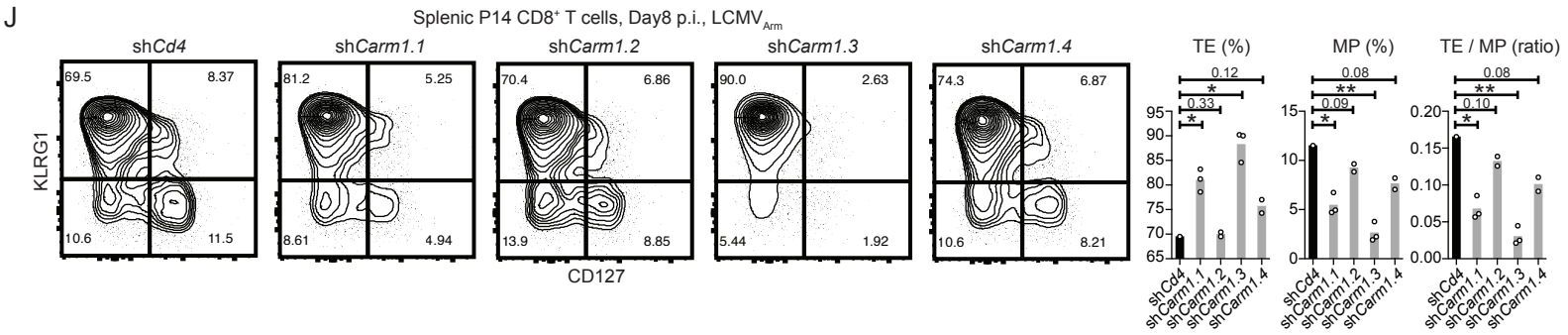
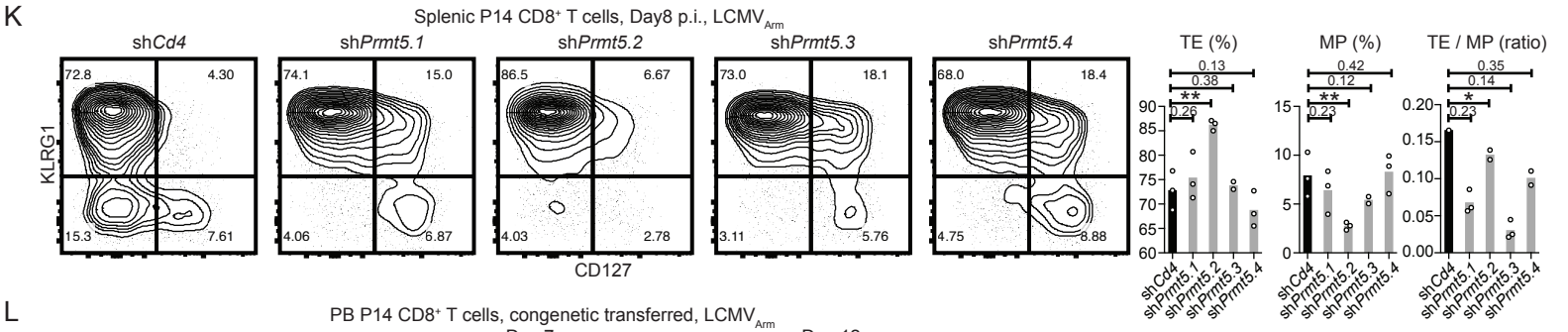

F
Input - Output
BAF
ISWI - Ino80
Chd
Histone Acetylation
Histone Methylation
Histone Ubiquitination
Arginine Methylation
DNA Methylation - Demethylation
HDAC
Histone Demethylation
Bromodomain
Chromodomain
G
EE - Other
H
MP - TE
I
K-Means clustering of CRF based on RNAi effect
MP v.s. TE
EE v.s. Other
Input v.s. Output
-4 -2 0 2 4

Figure S4


J
Splenic P14 CD8+ T cells, Day8 p.i., LCMVArm
shCd4
shCarm1.1
shCarm1.2
shCarm1.3
shCarm1.4
KLRG1
CD127
69.5
8.37
10.6
11.5
81.2
5.25
8.61
4.94
70.4
6.86
13.9
8.85
90.0
2.63
5.44
1.92
74.3
6.87
10.6
8.21
TE (%)
MP (%)
TE / MP (ratio)
0.12
0.33
0.08
0.09
0.08
0.10
shCd4
shCarm1.1
shCarm1.2
shCarm1.3
shCarm1.4


K
Splenic P14 CD8+ T cells, Day8 p.i., LCMVArm
shCd4
shPrmt5.1
shPrmt5.2
shPrmt5.3
shPrmt5.4
KLRG1
CD127
72.8
4.30
15.3
7.61
74.1
15.0
4.06
6.87
86.5
6.67
4.03
2.78
73.0
18.1
3.11
5.76
68.0
18.4
4.75
8.88
TE (%)
MP (%)
TE / MP (ratio)
0.13
0.38
0.26
0.42
0.12
0.23
0.35
0.14
0.23
shCd4
shPrmt5.1
shPrmt5.2
shPrmt5.3
shPrmt5.4

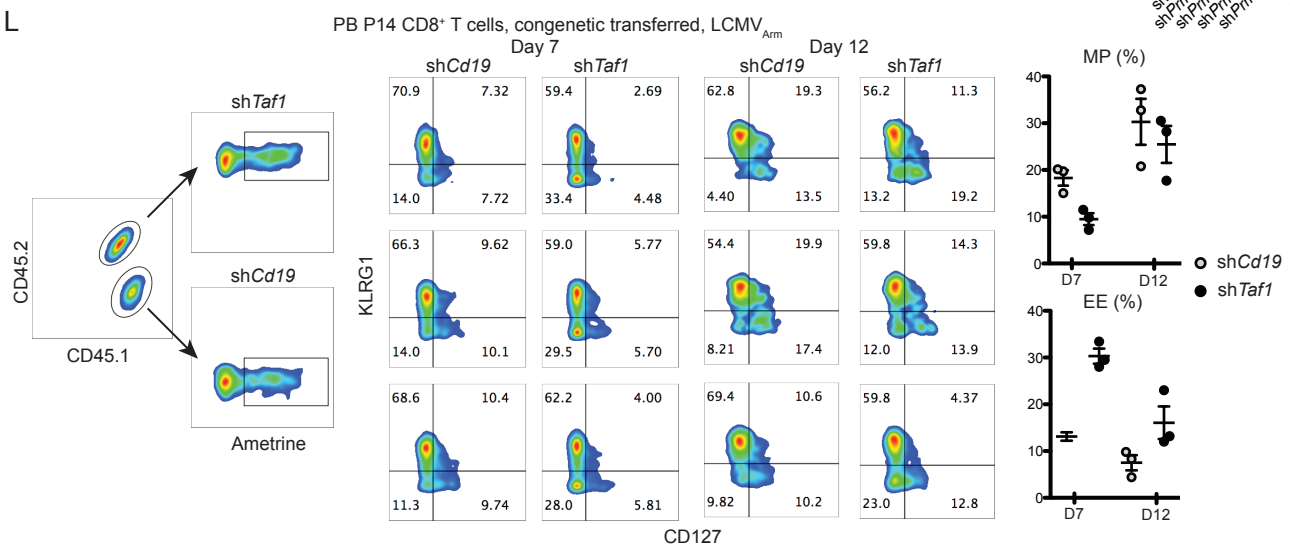

L
PB P14 CD8+ T cells, congenetic transferred, LCMVArm
Day 7
Day 12
shTaf1
shCd19
CD45.2
CD45.1
Ametrine
KLRG1
CD127
70.9
7.32
14.0
7.72
59.4
2.69
33.4
4.48
62.8
19.3
4.40
13.5
56.2
11.3
13.2
19.2
66.3
9.62
14.0
10.1
59.0
5.77
29.5
5.70
54.4
19.9
8.21
17.4
59.8
14.3
12.0
13.9
68.6
10.4
11.3
9.74
62.2
4.00
28.0
5.81
69.4
10.6
9.82
10.2
59.8
4.37
23.0
12.8
MP (%)
EE (%)
D7
D12
shCd19
shTaf1

Figure S4