Auto-aggressive CXCR6(+) CD8 T cells cause liver immune pathology in NASH

Citation for published version (APA):

Dudek, M., Pfister, D., Donakonda, S., Filpe, P., Schneider, A., Laschinger, M., Hartmann, D., Hueser, N., Meiser, P., Bayerl, F., Inverso, D., Wigger, J., Sebode, M., Oellinger, R., Rad, R., Hegenbarth, S., Anton, M., Guillot, A., Bowman, A., ... Knolle, P. A. (2021). Auto-aggressive CXCR6(+) CD8 T cells cause liver immune pathology in NASH. *Nature*, 592(7854). <https://doi.org/10.1038/s41586-021-03233-8>

Document status and date:

Published: 15/04/2021

DOI:

[10.1038/s41586-021-03233-8](https://doi.org/10.1038/s41586-021-03233-8)

Document Version:

Publisher's PDF, also known as Version of record

Document license:

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<https://doi.org/10.1038/s41586-021-03233-8>

Received: 17 February 2020

Accepted: 12 January 2021

Published online: 24 March 2021

Check for updates

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Nonalcoholic steatohepatitis (NASH) is a manifestation of systemic metabolic disease related to obesity, and causes liver disease and cancer^{1,2}. The accumulation of metabolites leads to cell stress and inflammation in the liver³, but mechanistic understandings of liver damage in NASH are incomplete. Here, using a preclinical mouse model that displays key features of human NASH (hereafter, NASH mice), we found an indispensable role for T cells in liver immunopathology. We detected the hepatic accumulation of CD8 T cells with phenotypes that combined tissue residency (CXCR6) with effector (granzyme) and exhaustion (PD1) characteristics. Liver CXCR6⁺ CD8 T cells were characterized by low activity of the FOXO1 transcription factor, and were abundant in NASH mice and in patients with NASH. Mechanistically, IL-15 induced FOXO1 downregulation and CXCR6 upregulation, which together rendered liver-resident CXCR6⁺ CD8 T cells susceptible to metabolic stimuli (including acetate and extracellular ATP) and collectively triggered auto-aggression. CXCR6⁺ CD8 T cells from the livers of NASH mice or of patients with NASH had similar transcriptional signatures, and showed auto-aggressive killing of cells in an MHC-class-I-independent fashion after signalling through P2X7 purinergic receptors. This killing by auto-aggressive CD8 T cells fundamentally differed from that by antigen-specific cells, which mechanistically distinguishes auto-aggressive and protective T cell immunity.

Preclinical mouse models—which reflect key findings in patients with NASH, including obesity, insulin resistance, metabolic disturbance, low-grade inflammation and liver cancer¹—are suitable for studying the pathogenesis of NASH. Mice fed a choline-deficient, high-fat diet (CD-HFD) or a Western diet for 9–12 months show these features of NASH, alongside with changes in the immune cell populations of the liver^{4–7}.

Liver CXCR6⁺ CD8 T cells in NASH

The development of NASH features in mice fed a CD-HFD requires conventional T cells, as shown by the fact that serum alanine aminotransferases and liver histology in mice that lack the genes that encode T cell receptor (TCR)β and TCRδ (*Tcrb*^{−/−} *Tcrd*^{−/−} mice) were normal (Fig. 1a–c),

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Fig. 1 | CD8 T cells in NASH mice have a unique pathogenic profile. a–c, Serum alanine aminotransferases (sALT) (**a**), nonalcoholic fatty liver disease (NAFLD) activity score (NAS) (**b**) and liver immunohistochemistry (**c**) in *Tcrb*^{−/−}*Tcrd*^{−/−} (*Tcrb*^{−/−} mice (*n* = 5) compared to wild-type (WT) littermates (*n* = 4) after six months of feeding a CD-HFD. Two independent experiments. Scale bar, 100 μ m. H&E, haematoxylin and eosin; ND, normal diet. **d**, Hepatic (left and middle) and splenic (right) CXCR6⁺CD44⁺ CD8 T cells. **e**, Frequencies of hepatic lymphocyte populations in mice fed a normal diet (*n* = 11) and NASH mice (*n* = 13). Two independent experiments. NK, natural killer cells; NKT, natural killer T cells. **f**, Phenograph-based analysis of hepatic lymphocytes (*n* = 5). *t*-SNE, *t*-distributed stochastic neighbour embedding. **g**, Representative

microscopic image of liver CD8 T cells depicting 3D cell objects. Scale bar, 50 μ m. **h–j**, Principal component (PC) analysis (**h**), volcano plot analysis (**i**) and GSEA (**j**) of differentially expressed genes from hepatic CD44⁺CXCR6⁺ CD8 T cells compared to CD44⁺CXCR6[−] CD8 T cells and splenic CD44⁺ CD8 T cells isolated from mice fed a normal diet (*n* = 3) or a CD-HFD (*n* = 3). Technical duplicates. FDR, false discovery rate. Exact *P* (**a**, **b**, **e**) and *n* (**e**) values are presented in Source Data. **P* < 0.05, ***P* < 0.01, ****P* < 0.001, *****P* < 0.0001; NS, not significant. One-way analysis of variance (ANOVA) with Tukey's (**a**, **b**), and two-way ANOVA with Sidak's (**e**), multiple comparison test. In **a**, **b**, **e** data are mean $\pm$ s.e.m., errors are shown as s.d.

whereas *Ja18*^{−/−} (which contain a knockout of the *Tcr*-*J* gene cluster) or *Cd1d*^{−/−} mice (*Cd1d* is also known as *Cd1d1*)—which lack T cells with invariant T cell receptors and natural killer T cells—developed features of NASH (Extended Data Fig. 1). Furthermore, *Tcrb*^{−/−}*Tcrd*^{−/−} mice fed a CD-HFD were not obese and liver steatosis was reduced (as compared to wild-type mice) (Fig. 1c, Extended Data Fig. 1), which collectively implicates T cells in the development of obesity, steatosis and NASH.

We phenotyped T cells, which revealed an increase of hepatic CD8 T cells in NASH mice that express the chemokine receptor CXCR6 (a marker of liver-resident CD8 T cells) but not CD49a, CX₃CR1 or KLRG1 (Fig. 1d, e, Extended Data Fig. 1), which characterize tissue-resident memory, effector memory or short-lived effector cells, respectively^{8–12}. High numbers of hepatic CXCR6⁺ CD8 T cells in NASH mice were distributed throughout the liver and co-expressed markers associated with exhaustion (PD1), effector function (GzmB, TNF and IFN γ) and apoptosis, but not Ki67 or BCL2 (which are markers of proliferation or survival) (Fig. 1f, g, Extended Data Fig. 1). An increase in the numbers of hepatic CXCR6⁺PD1^{high} CD8 T cells was a general feature of liver immune pathology driven by obesity, from a modest increase in the livers of mice fed a high-fat diet to a high increase in mice fed a Western diet or in obese, hyperphagic leptin-deficient *ob/ob* mice (Extended Data Fig. 2). The phenotype of hepatic CXCR6⁺ CD8 T cells we observed is not compatible with current classifications into liver-resident memory T cells or exhausted T cells^{8,13–15}, which led us to characterize these cells in more depth.

Our genome-wide transcriptome analysis revealed a clear separation of hepatic CXCR6⁺ from CXCR6[−] CD8 T cells in NASH mice, as compared

to CXCR6⁺ and CXCR6[−] CD8 T cells in the livers of healthy mice or splenic CD8 T cells (Fig. 1h). In liver CXCR6⁺ CD8 T cells from NASH mice, we found a higher expression of genes relevant for effector function (those encoding granzymes, as well as *Bhlhe40*) and exhaustion (*Pdcd1* and *Tox*), as well as the expression of genes indicative of tissue residency (for example, *Sitpr1*, *Klf2* and *Rgs1*) (Fig. 1i, Extended Data Fig. 3, Supplementary Table 1). We confirmed this using gene-set enrichment analysis (GSEA), which identified an enrichment of genes associated with T cell effector function and activation, exhaustion and tissue residency, without enrichment for genes dependent on T helper 17 cells or IL-17 (Fig. 1j, Extended Data Fig. 3), which have previously been associated with immune pathology during infection¹⁶. Consistent with a short half-life of hepatic CXCR6⁺ CD8 T cells in NASH, apoptosis-related (but not cell-cycle-related) genes were enriched (Fig. 1j, Supplementary Table 2). Together, this characterized hepatic CXCR6⁺ CD8 T cells in NASH as short-lived tissue-resident effector cells that receive concurrent activation and regulatory signals, and raised the question of how they were generated.

IL-15 induces hepatic CXCR6⁺ CD8 T cells

Our transcriptome profiling of hepatic CXCR6⁺ CD8 T cells did not show evidence for cardinal transcription factors that are required for the development of effector CD8 T cells, such as BLIMP1, EOMES or T-BET¹⁷ (Extended Data Fig. 3). Using ‘Binding Analysis for Regulation of Transcription’ (BART) program¹⁸, we identified 42 potentially relevant transcription factors, which—together with an in silico analysis that

Fig. 2 | High similarity between hepatic CXCR6⁺PD1^{high} CD8 T cells in NASH mice and patients with NASH. a, b, Correlation of hepatic CXCR6⁺CD8 T cells with sALT in NASH mice (**a**) and patients with NASH (compared with unaffected individuals) (**b**). **c**, Fold change in serum cytokine levels. **d**, Correlation of mouse CXCR6⁺PD1^{high} CD8 T cells with serum levels of IL-15. Four independent experiments. **e**, Hepatic expression of *IL15* expression in patients with NASH. **f**, Dot-plot heat map, showing expression and cellular detection rate of selected cluster marker genes in hepatic T cells of patients with steatosis (n = 4) or NASH (n = 3). MAIT, mucosal-associated invariant T cells. **g, h**, GSEA from differentially expressed genes of single-cell RNA-sequencing (RNA-seq) data from hepatic CD3⁺ T cells from patients with NASH, using gene sets for liver

residency³³ and IL-15 stimulation³⁴. **i**, Similarity of RNA-seq datasets from CXCR6⁺ compared to CXCR6⁺ CD8 T cells of NASH mice with human CD8 RGS1⁺ cells from patients with NASH. The solid line is the linear regression, and the grey area shows the 95% confidence intervals. **j, k**, Hepatic CXCR6⁺PD1^{high} CD8 T cells in NASH mice after anti-CD122 application and change in sALT. Exact *P* values (**e, j, k**) and *n* (**a–e, j, k**) values are presented in Source Data. **P* < 0.05, ***P* < 0.01. Coefficient of determination (*R*²) and statistical significance (*P* value) were determined using Pearson's (**a, b, d**) or Spearman's (**i**) correlation. One-way ANOVA with Tukey's multiple comparison test (**j**) and unpaired two-tailed *t*-test (**k**). In **e, j, k**, data are mean ± s.e.m., errors are shown as s.d.

revealed 21 downstream transcription factors—suggested a key role for the downregulation of FOXO1 in hepatic CXCR6⁺ CD8 T cells in NASH mice. GSEA confirmed the reduced expression of FOXO1-dependent genes, and reduced levels of FOXO1 protein in hepatic CXCR6⁺ CD8 T cells, but not in CD4 T cells, further corroborated these findings (Extended Data Fig. 3, Supplementary Tables 2, 3). To investigate the relevance of FOXO1 regulation, we performed loss- and gain-of-function experiments. Inhibition of FOXO1 in CXCR6⁺ CD8 T cells caused upregulation of CXCR6, whereas overexpression of FOXO1 in CD8 T cells reduced expression of CXCR6 (Extended Data Fig. 4). Accordingly, levels of FOXO1 in CD8 T cells inversely correlated with percentages of hepatic CXCR6⁺ CD8 T cells and with liver damage in livers of NASH mice and patients with NASH over the time of NASH development (Extended Data Fig. 4). Increased numbers of mouse and human hepatic CXCR6⁺ CD8 T cells, as well as increased human-liver CXCR6 expression and CXCR6 mRNA-expressing cells (Fig. 2b, Extended Data Fig. 4), demonstrated that similar hepatic CD8 T cell phenotypes correlated with liver damage in NASH mice and patients with NASH.

We screened for cytokines to identify mediators that downregulate FOXO1 in T cells, and detected increased levels of IL-15, IL-21 and TNF serum protein and liver mRNA in NASH mice (Fig. 2c, Extended Data Fig. 5). IL-15 signalling causes phosphorylation of FOXO1 for nuclear export and proteasomal degradation of cytosolic FOXO1^{19,20}, and has previously been associated with the development of NASH²¹. Upon treatment with IL-15 in vitro, mouse and human CD8 T cells downregulated

FOXO1 and upregulated CXCR6, PD1, Gzmb and CD69 (Extended Data Fig. 5). Of note, the expression of CXCR6 induced by IL-15 was further increased by IL-21 (Extended Data Fig. 5). Levels of serum IL-15 correlated with the frequency of hepatic CXCR6⁺PD1^{high} CD8 T cells, as well as the extent of liver damage and FOXO1 expression not only in NASH mice but also in patients with NASH (Fig. 2d, e, Extended Data Fig. 5), which together indicates a key role for IL-15 in NASH.

To further translate the findings from preclinical models into human disease, we isolated T cells from the livers of patients with steatosis or NASH and performed single-cell RNA sequencing (RNA-seq). We detected seven distinct hepatic CD8 T cell clusters, among which CD8 RGS1⁺ cells were increased in the liver of patients with NASH and co-expressed high amounts of CXCR6, PDCD1 and genes encoding granzymes and low levels of KLF2; in GSEA, this cluster of cells showed liver-residency and IL-15 signatures (Fig. 2f–h, Extended Data Fig. 6, Supplementary Tables 4–8). We also found RGS1⁺ CD8 cells in the fat tissue of patients with NASH (Extended Data Fig. 6, Supplementary Tables 9, 10). Importantly, the key gene signatures of hepatic CXCR6⁺ CD8 T cells from NASH mice were also present in hepatic RGS1⁺ CD8 T cells, but not in GNLV⁺FGFBP2⁺ CD8 T cells, from patients with NASH (Fig. 2i, Extended Data Fig. 6, Supplementary Table 11), which suggests similar functionality of CD8 T cells in NASH mice and patients with NASH. Upon exposure to IL-15, CD8 T cells downregulated CD122 (which is required for IL-15 signalling²²), consistent with CXCR6⁺, but not CXCR6⁺, CD8 T cells expressing CD122 (Extended Data Fig. 7); this suggested that

Fig. 3 | Acetate exposure enables CXCR6⁺ CD8 T cells to become auto-aggressive. **a**, Gzmb expression in IL-15-stimulated CD8 T cells exposed to metabolites ($n=3$). Untr., untreated. **b**, Liver acetate levels in mice fed a normal diet ($n=12$) and NASH mice ($n=13$). **c, d**, Auto-aggressive activity of IL-15-treated CXCR6⁺ CD8 T cells against hepatocytes in presence of acetate (15 mM) ($n=3$), or after transduction with pMP71-Foxo1-eGFP ($n=3$). **e, f**, Auto-aggressive activity of IL-15-treated, acetate-exposed CXCR6⁺ CD8 T cells in presence of anti-TNFR1 and anti-TNFR2 (anti-TNFR1/2) ($10 \mu\text{g ml}^{-1}$) or anti-IFN γ ($10 \mu\text{g ml}^{-1}$), or recombinant cytokines ($n=3$). **g**, sALT at day 2 after transfer of IL-15-exposed CD8 T cells and anti-TNF injection ($200 \mu\text{g per}$

mouse). Two independent experiments. **h–j**, Liver histology (**h**), NAS (**i**) and sALT (**j**) after anti-TNF application ($200 \mu\text{g per mouse twice a week, at 8 weeks}$) in mice fed a CD-HFD (normal diet, $n=11$; CD-HFD + immunoglobulin, $n=11$; CD-HFD + anti-TNF, $n=10$). Three independent experiments in **a, c–f**. MFI, mean fluorescence intensity. Exact P values (**a–g, i, j**) and n (**g**) values are presented in Source Data. * $P<0.05$, *** $P<0.001$, **** $P<0.0001$. Two-way ANOVA with Sidak's multiple comparison test (**a**), one-way ANOVA with Tukey's multiple comparison test (**c–g, i, j**) and unpaired two-tailed t -test (**b**). In **a–g, i, j**, data are mean $\pm$ s.e.m., errors are shown as s.d.

hepatic CXCR6⁺PD1^{high} CD8 T cells originated from CD122-expressing T cells. In NASH mice treated with anti-CD122 antibodies, only hepatic CXCR6⁺PD1^{high} CD8 T cells (and not CD4 T cells, CX₃CR1⁺ CD8 T cells or natural killer T cells) were depleted, and this treatment was followed by the amelioration of liver damage (Fig. 2j, k, Extended Data Fig. 7). Of note, the numbers of natural killer cells were also reduced after treatment with anti-CD122, but the number or phenotype of hepatic natural killer cells were not changed in NASH mice (Extended Data Fig. 7). These results suggested a role for IL-15 and CXCR6⁺ CD8 T cells in mediating liver damage in NASH.

Auto-aggression by CXCR6⁺ CD8 T cells

We next tested whether CXCR6⁺ CD8 T cells were pathogenic, but—despite high levels of Gzmb—they did not kill hepatocytes (Extended Data Fig. 8). Given the disturbance of lipid metabolism in the livers of patients with NASH², we evaluated the effect of fatty acids on T cell cytotoxicity. Only the short-chain fatty acid acetate further increased expression of Gzmb in CXCR6⁺ CD8 T cells and promoted the release of TNF, which is consistent with previous reports^{23,24} (Fig. 3a, Extended Data Fig. 8). Levels of hepatic acetate were elevated in NASH mice (Fig. 3b), which may be the result of gut dysbiosis²⁵ or of hepatocytes releasing acetate after exposure to fatty acids, and we found enrichment of acetate-induced—but not butyrate-induced—genes^{23,26} in hepatic CXCR6⁺ CD8 T cells from NASH mice (Extended Data Fig. 8, Supplementary Table 12). Notably, sequential application of IL-15 stimulation or FOXO1 inhibition followed by acetate exposure rendered CXCR6⁺FOXO1^{low} CD8 T cells capable of killing hepatocytes (Fig. 3d, Extended Data Fig. 8). We term this nonspecific killing activity of CXCR6⁺ CD8 T cells 'auto-aggression'. Overexpression of FOXO1 prevented auto-aggression by T cells (Fig. 3e, Extended Data Fig. 8),

which suggests that low FOXO1 activity rendered T cells responsive to environmental metabolic cues such as acetate. Adoptive transfer of either IL-15-stimulated or FOXO1-inhibited, and then acetate-exposed, CD8 T cells caused liver damage (Extended Data Fig. 8). This raised the question how acetate contributed to auto-aggression by CXCR6⁺ CD8 T cells.

Because acetate-exposed CXCR6⁺ CD8 T cells did not trigger exocytosis (which is required for the release of Gzmb from intracellular stores) and because *Prf1*^{−/−} mice (which lack perforin to deliver Gzmb-containing granules into target cells²⁷) developed NASH features (Extended Data Fig. 8), we analysed the effect of TNF on auto-aggression. Hepatocytes exposed to TNF upregulated ICAM1, which facilitated close interaction with LFA1-expressing CXCR6⁺ CD8 T cells; blockade of LFA1 prevented auto-aggression by T cells in vitro or after adoptive transfer in vivo (Extended Data Fig. 8). Consequently, TNF blockade prevented auto-aggression by T cells in vitro and in vivo, and hepatocyte incubation with TNF (but no other pro-inflammatory cytokines) sufficed to initiate auto-aggression by T cells even without prior exposure to acetate (Fig. 3e–g, Extended Data Fig. 8). Importantly, TNF inhibition for eight weeks in NASH mice reduced liver damage and hepatic steatosis (Fig. 3h–j), which together indicated a role for TNF in establishing physical contact between auto-aggressive CXCR6⁺ CD8 T cells and hepatocytes. In patients with NASH, we also found increased levels of *TNF* expression that correlated with liver damage (Fig. 3j, k), and this raised the question how pathogenic CD8 T cells were activated to exert auto-aggression.

Execution of CD8 T cell auto-aggression

MHC class I expression on hepatocytes was dispensable for auto-aggression by T cells, and anti-PD1 checkpoint inhibition—despite

Fig. 4 | Metabolic activation of CXCR6⁺ CD8 T cells for the execution of auto-aggression. **a**, Auto-aggression by IL-15-treated CXCR6⁺ CD8 T cells in the presence or absence of acetate against hepatocytes from wild-type or *B2m*^{-/-} mice, in presence of anti-MHC class I (10 μg ml⁻¹) or anti-PD1 antibodies (10 μg ml⁻¹) (*n* = 3). **b** CD8 T cell auto-aggression after inhibition of defined pathways (*n* = 3). **c**, Increase in intracellular calcium in auto-aggressive CXCR6⁺ CD8 T cells (*n* = 4). **d**, P2RX7 surface expression on CXCR6⁺ CD8 T cells (*n* = 7). **e**, Calcium influx in ATP-exposed CD8 T cells after P2RX7 inhibition (*n* = 8). **f, g**, Effect of P2RX7 or PANX1 inhibition (inh.) on auto-aggression and antigen-specific cytotoxic activity of IL-15-stimulated, acetate-exposed polyclonal or ovalbumin-specific OT1 CXCR6⁺ CD8 T cells (*n* = 3). **h**, Auto-aggression by IL-15 (15 ng ml⁻¹)-stimulated human CD8 T cells against allogeneic or autologous primary human hepatocytes (*n* = 3). **i**, Confocal

microscopic analysis and 3D reconstruction of cell surfaces in liver tissues from mice fed a CD-HFD or normal diet (green, CD8 T cells; red, cleaved caspase 3⁺ hepatocytes; grey, nonparenchymal cells); quantification in Extended Data Fig. 10 (*n* = 3). Scale bar, 30 μm. **j**, Auto-aggression of CXCR6⁺ CD8 T cells after anti-FasL in vitro (*n* = 3). **k**, Liver damage after anti-FasL application (100 μg per mouse twice a week, for 4 weeks) of mice fed a CD-HFD (normal diet, *n* = 11; CD-HFD + immunoglobulin, *n* = 11; CD-HFD + anti-FasL, *n* = 6). **l**, Model for CXCR6⁺ CD8 T cell auto-aggression in NASH. Three or more independent experiments in **a, b, d–g, j**. Exact *P* values (**a, b, d, e–g, j, k**) are presented in source data. **P* < 0.05, ****P* < 0.001, *****P* < 0.0001. Two-way ANOVA with Tukey's multiple comparison test (**e**), one-way ANOVA with Tukey's multiple comparison test (**a, b, f, g, j, k**) and unpaired two-tailed *t*-test (**b**). In **a, b, d, e–g, j, k**, data are mean ± s.e.m., errors are shown as s.d.

a high expression of PD1 by CXCR6⁺ CD8 T cells—did not augment auto-aggression (Fig. 4a). Consistent with this, neither phosphorylation of TCR-downstream signalling kinases nor immunometabolic features of effector function (such as a switch towards glycolysis²⁸) were observed in auto-aggressive CD8 T cells (Extended Data Fig. 9), which led us to search for other stimuli that triggered T cell activation in an MHC-independent fashion. Neither inhibition of phospholipase C nor mitogen-activated kinase-coupled signalling affected T cell auto-aggression, whereas inhibition of PI3K and calcium signalling abolished auto-aggression by CD8 T cells (Fig. 4b). Notably, incubation with ATP or nicotinamide adenine dinucleotide (NAD) (both of which are ligands for purinergic receptors²⁹)—but not with acetate, TNF, TLR ligands, chemokines or degradation products of ATP—triggered calcium influx into CXCR6⁺ CD8 T cells as a sign of activation (Fig. 4c, Extended Data Fig. 9). ATP was released through pannexin 1 (PANX1) (an oligomeric membrane channel complex that is responsible for releasing cytosolic ATP into the extracellular space^{29,30}) from dying hepatocytes or from hepatocytes cocultured with auto-aggressive CD8 T cells (Extended Data Fig. 9). Imaging using matrix-assisted laser desorption and ionization revealed an increased and uniform presence of fatty acids in the livers of NASH mice in contrast to the multifocal localization of ATP (Extended Data Fig. 9), which is compatible with ATP release at sites at which auto-aggressive CD8 T cells interacted with hepatocytes.

To investigate how extracellular ATP activated CD8 T cells to exert auto-aggression, we searched for purinergic receptors (such as

P2X7) that are expressed on tissue-resident CD8 T cells^{31,32}. We found P2RX7 expression on CXCR6⁺ CD8 T cells, which was required for auto-aggressive CXCR6⁺ CD8 T cells to respond to ATP or NAD and to supernatants from dying hepatocytes (Fig. 4d, e, Extended Data Fig. 9). Consequently, inhibition of P2RX7 or PANX1 prevented auto-aggression by CD8 T cells, whereas the antigen-specific cytotoxicity of CD8 T cells was not affected (Fig. 4f, g, Extended Data Fig. 9). Thus, distinct mechanisms discriminate auto-aggression, which is triggered by increased calcium influx through purinergic signalling via P2RX7 in auto-aggressive CXCR6⁺ CD8 T cells, from MHC-class-I-dependent cytotoxicity exerted by antigen-specific CD8 T cells.

Given the high similarity in gene signatures of CXCR6⁺ CD8 T cells in livers of NASH mice and patients with NASH, we examined whether similar mechanisms were involved in auto-aggression by human CD8 T cells. Human CD8 T cells stimulated with IL-15 also showed MHC-class-I-independent auto-aggression, but did not require acetate or TNF to become auto-aggressive (Fig. 4h, Extended Data Fig. 10). Similar to auto-aggression by CD8 T cells in mice, calcium influx was required to trigger auto-aggression by human CXCR6⁺ CD8 T cells (Extended Data Fig. 10).

We next characterized the mechanism by which CXCR6⁺ CD8 T cells executed auto-aggression. The exposure of CXCR6⁺ CD8 T cells to ATP led to rapid upregulation of FasL, but not TRAIL, and the death of hepatocytes depended on apoptosis, but not necroptosis; this was supported by the close contact between CD8 T cells and cleaved caspase 3⁺

hepatocytes in livers from NASH mice (Fig. 4i, Extended Data Fig. 10). Importantly, the blocking of FasL prevented auto-aggression by CD8 T cells in vitro and after adoptive transfer in vivo, and ameliorated liver damage in NASH mice (Fig. 4j, k, Extended Data Fig. 10).

Discussion

These results reveal that CD8 T cells are rendered auto-aggressive through defined, non-redundant sequential activation steps (Fig. 4l) and provide mechanistic insights into the pathogenesis of immune-mediated metabolic diseases such as NASH. The sequential steps for the generation of auto-aggressive mouse and human CD8 T cells involve IL-15-driven transcriptional programming followed by metabolic activation to execute auto-aggression. The auto-aggression of CD8 T cells in the liver may be involved in the development of liver cancer in patients with NASH that is driven by chronic liver damage, and may also be responsible for tissue damage in other organs. Our results demonstrate that auto-aggression by CD8 T cells is determined by mechanisms distinct from those of antigen-specific killing through CD8 T cells, which will help in the design of future immune therapies to avoid immune pathology without compromising the efficacy of antigen-specific T cell immunity.

Online content

Any methods, additional references, Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at <https://doi.org/10.1038/s41586-021-03233-8>.

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Methods

No statistical methods were used to predetermine sample size. The experiments were not randomized, and investigators were not blinded to allocation during experiments and outcome assessment.

Mice, diets and treatments

C57BL/6 mice were purchased from Charles River. H-2K^b-restricted TCR transgenic CD45.1⁺ OT-1 mice were bred and maintained under specific-pathogen-free conditions at the School of Medicine, TUM, according to the guidelines of the Federation of Laboratory Animal Science Association. Mice were housed with a light:dark cycle of 12 h, ambient temperature of 24 °C and humidity of 55%. Male mice between the ages of 6 and 10 weeks were used, and the number of mice subjected to experiments was chosen on the basis of previous studies^{4,5}. Six-week-old male C57BL/6, *Prf*^{-/-}, *Cd1d*^{-/-}, *Ja18*^{-/-} and *Tcrd*^{-/-} *Tcrg*^{-/-} mice were randomly fed ad libitum and housed at the German Cancer Research Center: normal diet, CD-HFD (Research Diets; D05010402), Western diet with *trans*-fat (Research Diets; D09100301, 40 kcal per cent fat (Primex shortening), 20 kcal per cent fructose, 2% cholesterol). For interventional studies, mice were fed a CD-HFD for the indicated time and randomly assigned to treatment groups with depleting monoclonal antibodies against CD122 (BioXcell, clone 5H4, no. BE0272, 100 µg per mouse), FasL (BioXcell, clone MFL3, no. BE0319, 100 µg per mouse) or TNF (BioXcell, clone XT3.11, no. BE0058, 200 µg per mouse) every 3 days. At the end of the experiment, mice were killed, and organs and serum were collected for analysis. All mice experiments were authorized by permission of the Regierung of Oberbayern (55.2-1-54-2532-60-14) and the animal research committee at the DKFZ (g129/16).

Human specimens

All experiments with human liver samples for in situ hybridization, isolating lymphocytes, RNA and measuring acetate concentrations were approved by the ethics committee of Hamburg (PV5184 and PV5187), Innsbruck (AN2017-0016 369/4.21) and Bern (2020-00852). Single-cell RNA-seq-analysis of CD3⁺ lymphocytes from human liver tissue and mesenteric fat tissue was approved by the Medical Ethics Committee of Heidelberg University (S-629/2013) and the Ethics Committee of Freiburg University (544/17). Experiments with liver samples from unaffected participants for isolating primary human hepatocytes were approved by the ethics committee at TUM (232/19S). Liver samples were acquired by mini-laparoscopy-guided liver biopsy or bariatric surgery. All studies were performed in accordance with the regulations of the ethics committee and the ethical guidelines of the World Medical Association (WMA) Declaration of Helsinki. The participants were recruited into the study after they gave written and informed consent.

Immune cell isolation

For the isolation of splenic immune cells, the spleen was removed and mechanically disrupted, filtered through a 70-µm strainer and red blood cells were lysed with a 2-min incubation with ammonium chloride potassium buffer. Spleen cells were washed twice with cold PBS. For isolation of hepatic immune cells, mouse liver was perfused via the portal vein for a minute and subsequently passed through a 100-µm mesh sieve, washed and the cell pellet was resuspended in 8 ml of GBSS containing collagenase IV (1:400, Worthington, no. LS004189) and digested in a rotatory shaker (240 rpm) for 15 min at 37 °C. The cell suspension was washed and lymphocytes were isolated through Percoll (40/80, GE Healthcare, no. 10607095) density centrifugation (7 acceleration, 1 deceleration, 1,440g, for 25 min). For stimulation of lymphocytes, cells were stimulated with IL-15–IL-15Rα and (10 ng ml⁻¹, ThermoFisher, no. 14-8152-80), IL-21 (10 ng ml⁻¹, Peprotech, no. 210-21), IL-7 (10 ng ml⁻¹, Peprotech, no. 217-17), IL-2 (10 ng ml⁻¹), TGFβ (10 ng ml⁻¹, Peprotech, no. 100-21), IFNγ (10 ng ml⁻¹, Peprotech, no. 315-05), IL-1β

(10 ng ml⁻¹, Peprotech, no. 211-11B) or TNF (5 ng ml⁻¹, ThermoFisher, no. PMC3013) for 18 h followed by flow cytometry analysis.

For isolation of lymphocytes from kidney, the renal capsula was removed and kidneys were digested by injecting 1 ml PBS containing type IV collagenase (1 mg ml⁻¹, Worthington, no. LS004189) and DNase I (50 U ml⁻¹, Roche, no. 04716728001). After mechanical dissection, tissue was incubated for 60 min at 37 °C, the cell suspension was filtered through a 70-µm mesh strainer, washed and subjected to Percoll (40%/80%) density centrifugation (1,440g, for 25 min). For isolation of lymphocytes from mouse skin, ears were cut and the dorsal side was removed from the ventral side. After cutting into small pieces, tissue suspensions were digested with PBS solution containing type IV collagenase (2 mg ml⁻¹) and DNase I (50 U ml⁻¹) for 60 min at 37 °C. Cell suspensions were filtered through 70-µm mesh, washed and used for analysis. For isolation of lung lymphocytes, both lobes were removed and digested by injecting 1 ml PBS solution containing type IV collagenase (1 mg ml⁻¹) and DNase I (50 U ml⁻¹). After tearing the lobes apart and removing remaining air by pipetting, the tissue suspension was digested for 30 min at 37 °C. Finally, the tissue suspension was filtered through a 70-µm mesh, washed and subjected to Percoll (40%/80%) density centrifugation (1,440g, 25 min).

To generate cell suspensions from human liver tissue, tissue was filtered through a 100-µm mesh and centrifuged (400g, 5 min). After cautiously decanting the supernatant, cell pellets were resuspended in medium (RPMI 1640 + 10% fetal calf serum (FCS) + 1% penicillin–streptomycin) and centrifuged before use in experiments.

Isolation of primary mouse hepatocytes

Mouse livers were perfused via the portal vein for 8 min with buffer containing 0.12 U ml⁻¹ collagenase (SERVA, no. 17465), were mechanically dissected, filtered through a mesh (300 µm) and centrifuged at 50g for 2 min to remove cellular debris. Hepatocytes were washed twice, filtered through a mesh (100 µm) and isolated by Percoll (80%/50%) gradient centrifugation (600g, 20 min). Freshly isolated primary mouse hepatocytes were seeded on collagen R (0.02%, SERVA, no. 47254.02)-coated 96-well E-plates (ACEA Biosciences, no. 2801035.6) in William's E medium (PAN Biotech, no. P04-29050) containing 200 mM glutamine (ThermoFisher, no. 25030081), 1 M HEPES pH 7.4, 10,000 U ml⁻¹ penicillin–streptomycin (Merck, no. P0781), 50 mg ml⁻¹ gentamycin (Merck, no. G1397), 0.005 ng ml⁻¹ insulin (INSUMAN rapid, Sanofi), 1.6% DMSO (Merck, no. 472301) and 10% FCS (PAN Biotech, no. P30-3306). After seeding, the medium was exchanged with a medium containing 1% FCS, and hepatocytes were maintained until their use in experimental approaches.

Isolation of primary human hepatocytes

Isolation of primary human hepatocytes was performed as previously described³⁵. In brief, liver tissue was perfused via 2–3 vessels at 37 °C in a two-step procedure using an EGTA-containing washing buffer followed by a digestion buffer containing 0.05% collagenase IV (Worthington). Viable hepatocytes were enriched using a Percoll gradient centrifugation and cell viability was determined by Trypan blue dye exclusion. Cells were seeded and cultivated on plates coated with rat tail collagen (collagen R, Serva) in Williams E medium containing insulin–transferin–selenium, 1.5% bovine serum albumin, GlutaMAX, non-essential amino acids, 100 U ml⁻¹ penicillin–streptomycin (all supplements from Gibco) in the presence of 10% FCS for 16 h. Hepatocytes were maintained in medium lacking FCS for 5 days before use in experiments.

Antibodies for flow cytometric analysis

For mouse studies, the following antibodies were purchased from Thermo Fisher or BioLegend, if not indicated differently: anti-BLIMP1 (5E7; 1:100), anti-BCL2 (100, 1:100), anti-BCL6 (BCL-DWN, 1:100), anti-CD3 (17A2, 1:250), anti-CD4 (PM4-5, 1:250), anti-CD8 (53-6.7, 1:250), anti-CD11a (1217, 1:250), anti-CD44 (IM7, 1:250, anti-CD45.1 (A20,

1:250), anti-CD49a (HMa1, 1:200), anti-CD54 (ICAM1, YN1/1.7.4, 1:250), anti-CD62L (MEL14, 1:250), anti-CD69 (HL2F3, 1:250), anti-CD107a (LAMP1, eBioD4B, 1:200), anti-CD122 (IL2rb, 5H4, 1:250), anti-CD178 (FasL, MFL3, 1:100), anti-CD183 (CXCR3-173, 1:250), anti-CD186 (CXCR6, SA051D1, 1:250), anti-CD223 (LAG3, eBioC98B7W, 1:250), anti-CD253 (TRAIL, N2B2, 1:200), anti-CD279 (PD1, 29F.1A12, 1:250), anti-CX3CR1 (SA011F11, 1:250), anti-EOMES (Dan11mag, 1:100), anti-FOXO1 (C29H4, Cell Signaling, no. 2880S, 1:200), anti-GzmB (GB11, 1:100), IFN γ (XM61.2, 1:100), anti-Ki67 (SolA15, 1:100), anti-KLRG1 (2F1, 1:200), anti-NK1.1 (PK136, 1:250), anti-P2XR7 (1F11, 1:100), anti-pAKT (SDRNR, 1:100), anti-pZAP70 (h3kobu5, 1:100), anti-pLCK (SRRCHA, 1:100), anti-T-BET (eBio4B10, 1:100), anti-TCR β (H57-597), anti-TNF (MP6-XT22, 1:100) and TOX (TXRX10, 1:100).

For human studies, the following antibodies were used: anti-CD3 (SK7), anti-CD8 (HIT8a), anti-CD45 (HI-30), anti-CD69 (FN50), anti-CD186 (K041E5), anti-GzmB (GB11), and anti-FOXO1 (C29H4).

Flow cytometric analysis of cell phenotype and function

Cell-surface staining with fluorochrome-labelled antibodies was performed for 30 min at 4 °C in PBS (ThermoFisher) supplemented with 2% FCS (Merck). Intracellular cytokine staining was performed using the Transcription Factor Staining Buffer Set (ThermoFisher, no. 00-5523-00) according to the manufacturer's instructions. In brief, liver-associated lymphocytes were incubated with PMA (20 ng ml⁻¹, Merck, no. P8139) and Ionomycin (500 ng ml⁻¹, Merck, no. I0634) for 4–5 h at 37 °C in the presence of brefeldin A (1:1,000, ThermoFisher, no. 00-4506-51) and monensin (1:1,000, ThermoFisher, no. 00-4505-51). Cells were labelled with Zombie NIR fixable viability kit (1:1,000, BioLegend, no. 423106) or with eBioscience fixable viability dye eFluor 780 (1:2,000, ThermoFisher, no. 65-0865-18). After cell-surface staining, cells were fixed, permeabilized and stained with fluorochrome-labelled antibodies against IFN γ , TNF and GzmB. Intracellular transcription factor staining was performed using the FOXP3/Transcription Factor Staining Buffer Set (no. 00-5523-00, ThermoFisher). For FOXO1 staining, cells were incubated with the unconjugated primary rabbit anti-mouse antibody overnight at 4 °C followed by incubation with an Alexa Fluor 488- (Cell Signaling, no. 4412s) or Alexa Fluor 647- (ThermoFisher, no. A-21244)-conjugated secondary goat anti-rabbit antibody for 1 h. For staining of phosphorylated proteins, cells were fixed and permeabilized in ice-cold methanol (100%). Flow cytometric analysis was performed using SP6800 (Sony); cells were sorted using SH800 (Sony). Data were analysed with FlowJo software (Tree Star, v.10.8).

To measure intracellular calcium concentration in T cells, the calcium-sensitive dye eF514 (2 mM, ThermoFisher, no. 65-0859-39) was used together with fluorochrome-labelled antibodies and cells were analysed by flow cytometry. Apoptosis in T cells was assessed by flow cytometry using APC- or FITC-labelled annexin V (Apoptosis Detection Kit, BioLegend: no. 640932 and no. 640914) following the manufacturer's instructions.

Phenographic analysis of hepatic T cells in mice fed a normal diet or CD-HFD for 12 months (normal diet, $n = 5$; CD-HFD, $n = 5$) was performed using single .fcs files of cells that were positive for TCR β and/or for NK1.1. Further analysis was executed with the R package Cytofit³⁶. The .fcs files were concatenated with the t -SNE algorithm. To cluster the populations, we used Rphenograph in Cytofit with default settings. Annotation of each cluster was performed using expression of the following markers: CD4, CD8, CXCR6, PD1 and GzmB.

Image-stream analysis

To visualize intracellular localization of FOXO1 protein, stained T cells were analysed using the Amnis ImageStream (Millipore). Similarity score between FOXO1 and propidium iodide (PI) (ThermoFisher, no. P1304MP) was calculated using the IDEAS software v.6.2. Similarity score was used to determine nuclear localization of FOXO1 and subsequent mean fluorescence intensity of the protein.

Stimulation of CD8 T cells in vitro

One million per ml splenic CD8 T-cells from C57BL/6 mice or from transgenic OT-1 mice were purified by immunomagnetic separation using AutoMACS (Miltenyi Biotec), were activated with plate-bound anti-CD3 (2 μ g ml⁻¹) in RPMI 1640 medium containing FCS (10%), L-glutamine (2 mM), penicillin–streptomycin (100 U ml⁻¹), β -mercaptoethanol (0.1 mM) and IL-2 (100 U ml⁻¹) for 2 days in 24-well plates. Cells were removed and transferred to a new plate at a concentration of 10⁶ cells per ml in the presence of TGF β (5 ng ml⁻¹) that was shown to be involved in T cell tissue residency³⁷. After 2 days, vital cells were isolated through Pancoll density centrifugation (1,440g, 20 min). Vital CD8 T cells were stimulated with IL-15–IL-15R α (10 ng ml⁻¹, ThermoFisher, no. 14-8152-80) and TGF β (5 ng ml⁻¹, Peprotech, no. 100-21) for 24 h to become highly enriched for CXCR6⁺, GzmB⁺ and CD69⁺ T cells. Positively selected human CD8 T cells were stimulated with IL-15 (15 ng ml⁻¹, Peprotech, no. 200-15) or TGF β (25 ng ml⁻¹, Peprotech, no. 100-21) as control for 5 days before analysis if not differently indicated. To be consistent in the terminology for IL-15 treatment of mouse and human T cells, we used the term IL-15 instead of IL-15–IL-15R α in the Article. For analysis of ligand-dependent influx of calcium in IL-15–IL-15R α -stimulated CXCR6⁺ CD8 T cells, the following molecules were used: LPS (100 ng ml⁻¹, Merck, no. L4524), CpG (10 μ g ml⁻¹, Invivogen, no. tlr1-1585), Poly I:C (10 μ g ml⁻¹, Invivogen, no. tlr1-picw), adenosine (1 μ M, Merck, no. A4036), ATP disodium salt hydrate (200 μ M, Merck, no. A26209), NAD (10 μ M, Merck, no. NAD100-RO) and recombinant CXCL16 (10 ng ml⁻¹, Peprotech, no. 250-28). For analysis of metabolite-dependent expression of GzmB in IL-15–IL-15R α -stimulated CXCR6⁺ CD8 T cells, the following substances were used (all from Merck): sodium lactate (5 mM, no. 71718), sodium citrate (5 mM, no. PHR1416), sodium pyruvate (5 mM, no. P5280), sodium acetate (15 mM, no. S2889), sodium propionate (0.5 mM, no. P1880), sodium butyrate (0.5 mM, no. 303410), palmitic acid (50 μ M, no. P0500), linoleic acid (100 μ M, no. L1012), oleic acid (100 μ M, no. O1383). Fatty acids were dissolved in ethanol. Medium pH values were monitored after adding these substances, but no change in pH value was detected.

Isolation of human CD8 T cells from peripheral mononuclear cells

CD8 T cells were generated from blood of healthy, voluntary donors after they gave written and informed consent. This study was approved by a vote from the ethics committee of the University Hospital München rechts der Isar. Blood was mixed with PBS (1:1) and loaded on Pancoll (PAN Biotech, no. P04-601000) to isolate peripheral blood mononuclear cells through centrifugation (1,400g, 25 min). Cells were washed, counted and mixed with human anti-CD8 magnetic beads (Miltenyi Biotec, no. 130-045-201) and subjected to immunomagnetic separation before use in experiments.

CD8 T cell cytotoxicity assay

Measurement of the dynamics of CD8 T cell cytotoxicity against primary mouse or human hepatocytes over time was performed with the impedance-based technology using an xCelligence RTCA MP device (ACEA Biosciences). In brief, after isolation, primary mouse or human hepatocytes were seeded in xCelligence plates (10⁴ cells per well) and on day 2 (after hepatocytes reached confluency and a maximum cell index) 3 $\times$ 10⁴ CD8 T cells were added. As indicated in the figures, acetate, recombinant cytokines, blocking antibodies or specific inhibitors were added. The following reagents were used: TNF (ThermoFisher, no. PMC3013), IL-6 (Miltenyi Biotec, no. 130-094-065), IL-1 β (Peprotech, no. 211-11B), pannexin I inhibitor (probenecid, Merck, no. P8761, 100 μ M), P2RX7 inhibitor (A-438079 hydrochloride, Cayman Chemical, no. 14580, 25 μ M), PLC γ inhibitor (edelfosine, Tocris, no. 3022, 10 μ M), EGTA (Merck, no. E3889, 2 mM), ERK inhibitor (U0126, Merck, no. 662005, 100 nM), PI3K inhibitor (wortmannin, Merck, no. W1628, 10 nM), calcineurin inhibitor (cyclosporin A, Merck, no. C3662,

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1 μM), anti-LFA1 (mouse: BioXcell, M17/4, BE0006, 10 $\mu\text{g ml}^{-1}$; human: BioLegend, 10 $\mu\text{g ml}^{-1}$), anti-mouse MHC1 (BioXcell, M1/42.3.9.8, no. BE0077, 10 $\mu\text{g ml}^{-1}$), anti-mouse PD1 (BioXcell, RMP1-14, no. BE0146, 10 $\mu\text{g ml}^{-1}$), anti-mouse TNFR1 (BioLegend, 55R-593, no. 113102, 10 $\mu\text{g ml}^{-1}$), anti-mouse TNFR2 (BioLegend, TR75-32.4, no. 113202, 10 $\mu\text{g ml}^{-1}$), anti-human TNF (BioLegend, MAb11, no. 502901), anti-mouse IFN γ (BioLegend, XMG1.2, no. 505801, 10 $\mu\text{g ml}^{-1}$), anti-FasL (mouse: BioXcell, MFL3, no. BE0319, 10 $\mu\text{g ml}^{-1}$; human: BioLegend, NOK-1, no. 306415, 10 $\mu\text{g ml}^{-1}$) and anti-human NKG2D (BioLegend, 1D11, no. 320813, 10 $\mu\text{g ml}^{-1}$). Hepatocytes treated with respective reagents alone were used for measurement of baseline values. Measured electrical impedance is shown as cell index and normalized to the time point of starting the coculture of T cells with hepatocytes. The cell index at 24 h after start of coculture was used to calculate the efficacy of hepatocyte killing. Percentage of killing was calculated by subtracting the cell index values from controls (hepatocytes alone or with respective substances).

Cytotoxicity measurement of human CD8 T cells against human K562 cells (ATCC, CCL-243) was performed by labelling target cells with CFSE (1 μM , ThermoFisher, no. C34554) and incubating with effector cells in different effector-to-target ratios for 24 h. Dead K562 cells in coculture with CD8 T cells were detected by flow cytometry and percentage was calculated to K562 cells alone.

Adoptive transfer of auto-aggressive CD8 T cells into C57BL/6 mice

CXCR6⁺ CD8 T cells from C57BL/6 mice were differentiated and stimulated with IL-15–IL-15R α (10 ng ml^{-1}) as described in ‘Stimulation of CD8 T cells in vitro’, followed by exposure to sodium acetate (15 mM, Merck) for 24 h. CD8 T cells were washed and adoptively transferred (4×10^6 cells per mouse) into mice in combination with antibodies against either TNF (200 μg per mouse, BioXcell, clone XT3.11, no. BE0058), LFA1 (100 μg per mouse, BioXcell, clone M17/4, no. BE0006), FasL (100 μg per mouse, BioXcell, clone MFL3, no. BE0319) or P2XR7 (50 μg per mouse, 13A7). sALT was determined two days after transfer.

Extracellular flux analysis

Extracellular acidification rates were measured in XF medium (no. 03334-100, Agilent, non-buffered RPMI 1640 containing 2 mM L-glutamine, and 1 mM sodium pyruvate) under basal conditions and in response to 10 mM glucose (Merck, no. G7021), 2 μM oligomycin (Merck, no. 75351) and 50 mM 2-deoxy-D-glucose (Merck, no. D8375) using a 96-well XF Extracellular Flux Analyzer with Analyzer Software v.1.4.2 (Agilent).

Retroviral infection and overexpression of FOXO1

Mouse FOXO1–eGFP was amplified from a GFP-Foxo1 plasmid using the following primers Foxo1 fw_SLIC: 5' CTCCAAGCTCACTTACAGGC GAGTCAACATGGCCGAGCG3'; eGFP rev_SLIC: 5' GCGGTAAGATGCTCG AATTCTTTACTTGTA CAGCTCGTCC 3' (underlining denotes restriction enzyme sites for KpnI and EcoRI). GFP-Foxo1 was a gift from D. Accilli (Addgene plasmid no. 17551; <https://www.addgene.org/17551/>; RRIDAddgene_17551). Foxo1–eGFP was inserted into pMP71 retroviral vector by sequence and ligation independent cloning³⁸ to generate pMP71-Foxo1–eGFP. Specifically, the eGFP of pMP71–eGFP was deleted by NotI + EcoRI restriction enzyme digestion and the PCR fragment was inserted. pMP71-Foxo1–GFP and control pMP71–eGFP containing only GFP were used for retroviral transduction of activated CD8 T cells: on day 1, the retroviral packaging cell line Plat-Eco (Cell Biolabs, no. RV-101) was transfected using Lipofectamine 3000 (ThermoFisher, no. L3000001) according to the manufacturer's instructions. On the same day, splenic CD8 T cells from C57BL/6J mice were isolated and stimulated with plate-bound anti-CD3 and anti-CD28 antibodies (5 ng ml^{-1}) in RPMI 1640 medium containing FCS (10%), L-glutamine (2 mM), penicillin–streptomycin (100 U ml^{-1}), β -mercaptoethanol (0.1 mM) and IL-2 (300 U ml^{-1}). On day 3, viral supernatant was

transferred to retronectin-coated 12-well plates (20 ng ml^{-1} , TaKaRa, no. T100A/B), and spun at 1,000g for 120 min at 37 °C. Activated CD8 T cells were loaded on centrifuged viral supernatant and this process was repeated on the next day. Transduced T cells were cultured for two days, adding fresh medium with IL-2 (300 U ml^{-1}) on day 1 after transduction. CXCR6, FOXO1 and GzmB expression were determined in GFP⁺ CD8 T cells, allowing us to differentiate between T cells that overexpressed FOXO1.

For CD8 T cell cytotoxicity experiments against primary mouse hepatocytes with Xcelligence device, transduced GFP⁺ CD8 T cells were sorted by fluorescence-activated cell sorting on day 1 after transduction. On day 2 after transduction, sorted and transduced CD8 T cells were stimulated with IL-15–IL-15R α (10 ng ml^{-1}) and exposed to acetate (15 mM) overnight and added to primary hepatocytes at an effector-to-target ratio of 3:1. After 24 h of coculture, auto-aggression was calculated as described in ‘CD8 T cell cytotoxicity assay’.

Measurement of serum parameters

Serum was obtained by cardiac puncture after mice were killed, and sALT levels were measured with the Reflotronplus system (Roche). Detection of cytokine concentrations in serum was performed with a 23-plex bead array (Bio-Rad, no. M60009RDPD) or an enzyme-linked immunosorbent assay for IL-15 (Peprotech, no. 900-K188) or IL-21 (Peprotech, no. 900-K368).

Fluorescent-linked immunospot (fluorospot) assay

One hundred thousand in vitro-differentiated CXCR6⁺ CD8 T cells were incubated for 24 h in the presence of TGF β (5 ng ml^{-1}), IL-15–IL-15R α (10 ng ml^{-1}) and sodium acetate (15 mM) on TNF- and IFN γ -coated 96-well plates. For antigen-specific stimulation, medium was supplemented with 1 nM SIINFEKL for the last four hours followed by detection of TNF- and IFN γ -secreting CXCR6⁺ CD8 T cells, according to the manufacturer's instructions (Immunospot, no. mT2001F, no. mT02, no. mT11). Counting of TNF- and IFN γ -producing CXCR6⁺ CD8 T cells was performed with Immunospot's S6 ULTIMATE Analyzer and Immunospot Software (CTL Immunospot 5.4 Professional DC Software).

Confocal microscopy of primary cells in vitro

Thirty thousand hepatocytes were seeded on collagen-coated sterile 12-well silicone chamber slides (Ibidi, no. 81201) and 1×10^5 CXCR6⁺ CD8 T cells were added with medium containing TGF β (5 ng ml^{-1}), IL-15–IL-15R α (10 ng ml^{-1}) and sodium acetate (15 mM). After 18 h of coculture, cells were fixed with 4% PFA, washed with PBS and were stained with ICAM1–PE (1:200), LFA1–Alexa Fluor 647 (1:200) and DAPI (1:2,000, Merck, no. 10236276001). Stained sections were mounted in Mowiol and analysed on an inverted TCS SP8 confocal microscope (Leica) for tiled imaging using a HC PL APO CS2 40 $\times$ /1.30 NA or a HC PL APO CORR CS2 63 $\times$ /1.30 NA objective. Tiled x–y stacks were acquired with 0.5- μm z-spacing to provide 3D image volumes of 7–11- μm depth. Confocal image analysis was performed using Imaris software (v.9.5, Bitplane) on maximum projections of 10 large z-plane sections. Automated analyses using the IMARIS surface generation tool were used to reconstruct surface objects for T cells and hepatocytes based on LFA1 and ICAM1 staining. Distances between surface objects were calculated on the basis of positional information using the IMARIS Shortest Distance Statistics tool.

Confocal microscopy and 3D reconstruction of liver tissue

Thirty-micrometre paraffin sections were cut on a HM355S microtome (ThermoFisher) and adhered to Superfrost Plus slides (ThermoFisher). Antigen retrieval was performed in 1 mM EDTA pH 9.0.05% Tween-20 at 100 °C for 20 min. Sections were then permeabilized and blocked in PBS containing 0.3% Triton X-100 (Sigma-Aldrich) and 10% FBS followed by staining in the same blocking buffer. The following primary antibodies were used for staining: rat anti-mouse CD8 (4SM15, ThermoFisher),

rabbit anti-mouse cleaved caspase 3 (D3E9, Cell Signaling Technology), goat anti-mouse collagen IV- $\alpha 1$ (polyclonal, Novus Biologicals). The following secondary antibodies were used for staining: Alexa Fluor 488 chicken anti-rat IgG (H+L), Alexa Fluor 568 chicken anti-rabbit IgG (H+L) and Alexa Fluor 647 donkey anti-goat IgG (H+L), (ThermoFisher).

Stained slides were mounted with Fluorsave (Merck Millipore) and images were acquired on an inverted Leica microscope (TCS STED CW SP8, Leica Microsystems) with a motorized stage for tiled imaging. To minimize fluorophore spectral spill over, we used the Leica sequential laser excitation and detection modality. The bleed-through among sequential fluorophore emission was removed by applying simple compensation correction algorithms to the acquired images. The semiautomatic surface-rendering module in IMARIS (Bitplane) was used to create 3D volumetric surface objects corresponding either to individual cells or to the liver vascular system. Signal thresholds were determined using the IMARIS Surface Creation module, which provides automatic threshold identification and value-based visual surface thresholds around the positively stained objects. The semiautomatic channel colocalization module in Imaris (Bitplane) was used to detect and reconstruct surface contact areas between CD8 T cells and cleaved-caspase-3-positive hepatocytes.

For confocal imaging of CXCR6 in the liver, livers were perfused with 2.5 ml Antigenfix solution (Diapath) via the portal vein, excised and fixed for 4 h in 1 ml Antigenfix. Fixed liver lobes were embedded in Tissue-Tek O.C.T. compound (Sakura Finetek) and frozen at -80°C before 50- μm cryosections were cut. Liver sections were permeabilized and blocked with 0.1 M Tris (AppliChem) containing 1% BSA, 0.3% Triton X-100 (Gebro Biotechnik), 1% normal mouse serum (Sigma) for 2 h. Sections were stained in blocking buffer with anti-CD3 (Biolegend, clone 17A2), anti-CD8 (eBioscience, clone 53-6.7), anti-CXCR6 (Sony, clone SA051D1), anti-PD-1 (R&D Systems, polyclonal) and anti-collagen IV (Abcam, polyclonal). Unconjugated primary antibodies were stained with the following secondary antibodies overnight: donkey anti-rabbit IgG (Biolegend, clone Poly4064) and donkey anti-goat IgG (Jackson Immunology, polyclonal). Stained sections were mounted in Mowiol and analysed on an inverted TCS SP8 confocal microscope (Leica) for tiled imaging using a HC PL APO CS2 20 $\times$ /0.75 IMM objective. Tiled x - y stacks were acquired with 1- μm z -spacing to provide 3D image volumes of 15- μm depth. Three-dimensional image analysis was performed using IMARIS 9.5 software (Bitplane) following adaptive image deconvolution using the Leica TCS SP8 LIGHTNING tool. The IMARIS surface generation tool was used to reconstruct 3D surfaces for CD3 $^{+}$ CD8 T cells, T cell subsets were identified based on mean voxel intensities for CXCR6 and PD1.

Multiplex immunohistochemistry

Sequential immunostaining was performed on 2- μm -thick formalin-fixed paraffin-embedded (FFPE) mouse liver sections, as previously described⁵. Heat-mediated antigen retrieval was performed in citrate buffer (ThermoFisher). Antibody elution was performed between each staining cycle using a 2-mercaptoethanol, SDS buffer. Primary antibodies used: anti-PD1 (1/200, R&D Systems, no. AF1021), anti-CD8a (1/200, ThermoFisher no. 14-0808-82), anti- α SMA (1/200, Agilent no. M085129-2), anti-IBA1 (1/1,000, EMD Millipore no. MABN92) and anti-CK19 (1/200, DSHB no. TROMA-III). TROMA-III was deposited to the DSHB by R. Kemler (DSHB Hybridoma Product TROMA-III).

Mass spectrometry imaging

N-(1-Naphthyl) ethylenediamine dihydrochloride (NEDC) (>98% purity, no. 222488) was purchased from Sigma Aldrich and used without further purification. Methanol (ULC-MS grade, no. 136841), water (LC-MS grade, no. 232178), ethanol (ULC-MS grade, no. 052541), acetone (LC-MS grade, no. 010378) and xylene (AR grade, no. 242505) were purchased from Biosolve. Haematoxylin (Merck, no. 104302) and eosin Y (J. T. Baker, no. L088-03) were used under standard laboratory protocols. Entellan was purchased from Merck (no. 107961). Liver tissue

was obtained from mice fed a normal diet or CD-HFD for 12 months and sections of 12- μm thickness were generated at -15°C with a cryomicrotome (Leica). Samples were thawed and mounted onto clean Intellislide (Bruker). Slides were then spray-coated with matrix using an HTX TM-Sprayer (HTX Technologies) under the following conditions: 7.5 mg ml^{-1} NEDC in ethanol/methanol/acetone, (50/49/1) spray temperature 80°C , spray height 40 mm, velocity 1,200 mm min^{-1} , flow rate 0.11 ml min^{-1} , gas flow rate 3 l min^{-1} , 3 mm track spacing, 12 passes. Final concentration of NEDC was approximately 2.75 $\mu\text{g per mm}^2$. Samples were analysed immediately after spray-coating, using a TimsTOF Flex mass spectrometer (Bruker). Mass spectrometry images (MSI) were generated for both positive and negative ionization modes, with the m/z window tuned to either 50–150 m/z (metabolites) or 70–900 m/z (lipids), with the pixel size set to 10 μm . All samples were analysed using the post-ionization functionality of the TimsTOF Flex. Individual pixels were gathered with 25 laser shots, with post-ionization laser delay of 9 μs . TimsTOF data files were imported into SciLS Lab (Bruker Daltonics). Post-MSI, samples were washed of matrix and H&E-stained according to standard laboratory protocols. Slides were imaged using an Aperio CS2 (Leica). Box plots of segmented liver tissues were generated, with root mean square normalization between liver tissue from mice fed a normal diet or CD-HFD.

Measurement of acetate

Mouse and human mouse tissues were weighed, 50 μl PBS was added and they were then gently centrifuged at 300g for 8 min. Organ supernatants were recovered and acetate contents determined using the acetate assay kit (no. MAK086, Merck) according to the manufacturer's instructions. Organ acetate levels were normalized to organ weight. Concentration of acetate in the supernatant of primary mouse hepatocytes that were treated with palmitate (250 μM) or with trimeric FasL (no. ALX-522-001-C010, Enzo) for 24 h was determined according to the manufacturer's instructions (no. MAK086, Merck).

Measurement of extracellular ATP

Supernatant of primary mouse hepatocytes that were treated with palmitate or trimeric FasL for 24 h or of CXCR6 $^{+}$ CD8 T cells were collected in vitro and the concentration of ATP was measured according to manufacturer's instructions (no. A22066, Thermo Fisher).

Histology, immunohistochemistry, scanning and automated analysis

Tissues were fixed in 4% paraformaldehyde, paraffin-embedded and treated as previously reported³⁹. Two- μm sections from FFPE or cryopreserved tissues were stained with haematoxylin and eosin or stained with specific antibodies (anti-CXCR6, 1:200, clone PA5-33462, Invitrogen; anti-ICAM1:1500, clone 3E2B, Linaris; anti-MHC I:1500, clone ER-HR 52) using a Leica Biosystems Bond MAX. Stained tissue slides were scanned using an Aperio System and analysed using Aperio Image Scope software 12.4.0. NAS was performed in a blinded fashion on mouse livers and cross-validated by an expert veterinary pathologist.

For in situ hybridization, freshly non-baked 5- μm FFPE tissue slices were cut and stained according to manufacturer's (ACD biotech) protocol for manual-assay RNAscope, using probes against human CXCR6 (468461) and mouse CXCR6 (no. 871991). For quantification of cells that were positive for CXCR6 mRNA per mm^2 liver tissue, positive cells of four independent regions of interests in the represented areas were counted and the average number was calculated.

Quantitative real-time PCR

RNA from liver tissue from mice fed a normal diet or CD-HFD for 12 months was isolated according to the manufacturer's instructions (no. T2010S, NEB). cDNA was generated using SensiFAST cDNA synthesis kit from Bioline (no. BIO-65054). Primers against mouse *Il15*, *Il15ra*, *Il21*, *Cxcl16*, *Tnf*, *Il6* and *Il1b* were purchased from Eurofins (Munich);

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the sequences are listed in the Supplementary Information. A mix of primer pairs, double-distilled water and 2× Takyon Mix SYBR green assay (Eurogentec) was added to the cDNA and amplified copies were quantified by LightCycler 480 (Roche).

RNA from human liver tissue was isolated according to the manufacturer's instructions (no. 740955.250, Macherey Nagel). cDNA was generated using high-capacity cDNA Reverse transcription Kit from Applied Biosystems (no. 4368813). Primers against *IL4* (no. Hs00174122_m1), *IL15* (no. Hs01003716_m1), *IL15RA* (no. Hs00542602_g1), *IL21* (no. Hs00222327_m1), *CXCR6* (Hs01890898_s1), *CXCL16* (Hs00222859_m1), *ICAM1* (no. Hs00222859_m1), *TNF* (no. Hs01113624_g1) and *HRPT1* (no. Hs02800695_m1) were purchased from Thermo Fisher Scientific. A mix of primer pairs, double distilled water and KAPA Probe Fast Universal (no. KK4715, KAP Biosystems) was added to the cDNA and amplified copies were quantified by ViiA7 (Applied Biosystems).

RNA-seq analysis

Samples for RNA-seq were directly sorted in a semi-skirted 96-well PCR plate including lysis buffer (2× TCL buffer (Qiagen, no. 1070498) + RNase-free water + β-mercaptoethanol (1% total volume)). Library preparation for bulk 3'-sequencing of poly(A)-RNA was performed as previously described⁴⁰. The barcoded cDNA of each sample was produced with a Maxima RT polymerase (Thermo Fisher) exercising oligo-dT primer containing barcodes, unique molecular identifiers (UMIs) and an adaptor 5' ends of the cDNAs were prolonged by a template switch oligo (TSO), and after uniting of all samples full-length cDNA was amplified with primers binding to the TSO site and the adaptor. cDNA was supplemented with the Nextera XT kit (Illumina) and 3'-end-fragments finally augmented with P5 and P7 Illumina overhangs. We sequenced the library using NextSeq 500 (Illumina). To generate the UMI tables for samples and genes we used Drop-seq pipeline (<https://github.com/broadinstitute/Drop-seq>). We used GRCh38 reference genome for the alignment and ENSEMBL annotation release 75 for annotating genes. The DESeq2 R package⁴¹ (v.2.1.28.1) was used to detect the differentially expressed genes (fold change 1.5 and adjusted *P* value ≤ 0.05). Principal component analysis was performed using the 'prcomp' function in R and visualized via ggplot R package (v.3.3.2). Volcano plots and heat maps were plotted using EnhancedVolcano and pheatmap R packages (v.1.0.12).

Identification of transcription factors and network analysis

Transcription factor analysis was performed using the differentially expressed genes in datasets from CXCR6⁺ and CXCR6⁻ mice fed a CD-HFD. To predict the upstream transcription factors that control the differentially expressed genes, we used the 'Binding Analysis for Regulation of Transcription' prediction tool (BART v.1.1)¹⁸ with default settings. We considered a transcription factor as significant at *P* value < 0.01. Transcription factors perturbed within the transcriptome datasets (downstream transcription factors) were identified using the mouse transcription factor atlas⁴². This analysis identified 25 transcription factors in the datasets from CXCR6⁺ versus CXCR6⁻ mice fed a CD-HFD. We generated transcription factor–transcription factor network of both upstream transcription factors from BART analysis and downstream transcription factors from the transcriptomes as follows: first, promoter sequences (−2 kb) of significantly differentially expressed genes from the datasets were downloaded from eukaryotic promoter database⁴³. Second, we retrieved the transcription factor binding sites from the JASPAR core and Hocomoco databases^{44,45}. Then, we scanned promoter sequences (−2-kb promoters) of differentially expressed genes and transcription factors for binding sites using a custom Python script. We generated transcription factor–transcription factor networks and visualized them in Cytoscape v.3.7.1 (ref. ⁴⁶). To assess the hierarchy of transcription factor networks, in (*I*) and out (*O*) degrees were computed for each transcription factor and its targets using the igraph R package (<https://igraph.org/>, v.1.2.6) and hierarchy height $H = (O - I) / (O + I)$ was

calculated as previously described⁴⁷. Hierarchy height score defines three levels of transcription-factor network hierarchy.

Human single-cell RNA-seq in patients with NASH

Sample generation. Single-cell suspensions from human liver were generated by cutting freshly obtained liver tissues into small pieces, followed by enzymatic digestion using the Miltenyi tumour dissociation kit (no. 130-095-929) according to the manufacturer's instructions. Liver pieces were digested for 20 min at 37 °C while shaking, and stopped by adding medium (RPMI + 20% FBS). Digested tissues were passed through a 70-μm cell strainer and centrifuged at 350g for 10 min. Red blood cell lysis was carried out with ACK lysis buffer (Thermo Fisher Scientific, no. A1049201) and, after washing, the remaining cells with complete medium were strained through a 45-μm mesh. CD45⁺ cells were sorted and subjected to single-cell RNA-seq analysis. Liver cells were stained with anti-CD45-PE (HI30, no. 12-0459-42) and Live/Dead discrimination (1:1,000, Thermofisher, no. L34973), washed with PBS + 0.05 mM EDTA (10 min, 300g, 4 °C) and sorted into single CD45⁺ cells using a FACSria FUSION. Library generation was performed according to the manufacturer's protocol (Chromium Next EM Single Cell 3' GEM, no. 10000128). Single-cell sequencing was performed with an Illumina NovaSeq-6000.

Single-cell RNA-seq analysis. To perform downstream analysis, we used filtered gene-barcode matrices generated from the 10X Genomics pipeline cellranger count to serve as input for the R package Seurat 3^{48,49} (v.3.2.2). Analysis was initiated by creating Seurat objects from liver samples from patients with steatosis (*n* = 4) or NASH (*n* = 3). We excluded genes that were expressed in fewer than three cells, and also cells with fewer than 200 genes. Datasets were normalized and cell-cycle regression was performed. We executed unsupervised clustering by computing principal components using the RUNPCA function in the Seurat R package. Cluster analysis was performed using the cluster R package⁵⁰ with a clustering resolution of 1, and cell clusters were visualized using uniform manifold approximation and projection (UMAP). Markers were identified using the 'Finding All Markers' function (logfc. threshold = 0.25 and Wilcoxon rank sum test-adjusted *P* value < 0.05). Dot plot and feature plots were visualized using the Seurat R package.

To compare mouse bulk RNA-seq dataset (CXCR6⁺ versus CXCR6⁻ mice fed a CD-HFD) with human single-cell RNA data from livers of patients with NASH (CD8 *RGS1*⁺ and CD8 *GNLY*⁺*FGFBP2*⁺) we converted mouse genes to human orthologues using an online tool (<https://biobdnet-abcc.ncifcrf.gov/db/dbOrtho.php>) and computed the Spearman correlation between the log₂-transformed fold changes of both datasets. Correlation was considered significant at *P* < 0.05, and visualized as a scatter plot using the ggplot2 package (v.0.4). To assess the similarity between CD8 T cell populations from fat tissue (CD8 *RGS1*⁺ and CD8 *CCL4*⁺) and livers of patients with NASH (CD8 *RGS1*⁺ and CD8 *PLCG2*⁺) single-cell datasets, we calculated Jaccard similarity co-efficient, $J = |P_i \cap P_j| / |P_i \cup P_j|$. *P_i* and *P_j* denote genes from the populations. The degree of similarity was visualized as a heat map using yjr pheatmap R package (v.1.0.12) (<https://cran.r-project.org/web/packages/pheatmap/index.html>).

GSEA

To execute GSEA, we downloaded the following gene sets for defined cells or settings in .gmt format from the molecular signature database (MsigDB) (<http://software.broadinstitute.org/gsea/msigdb/index.jsp>): (1) effector gene set (KAECH_NAIVE_VS_DAY8_EFF_CD8_TCELL_DN); (2) GPCR signalling pathway; (3) TCR signalling pathway; (4) T_H17 gene set; (5) cell cycle; (6) GSE46025_WT_VS_FOXO1^{-/-}_KLRG1_LOW_CD8_EFFECTOR_TCELL up and down gene sets; and (7) apoptosis hallmark gene set. To convert human gene sets into mouse orthologues, we used a shell script (<http://crazyhotommy.blogspot.com/2018/02/convert-human-gmt-file-to-mouse-for-gsea.html>) and compared these mouse genes with our RNA-seq dataset. Raw microarray data for the

transcription factors BATF and TBX21 were from the Gene Expression Omnibus (GEO) database (accession identifiers GSE54215 (BATF knockout versus wild type (day 3)) and GSE122931 (TBX21 knockout versus wild type (H1 cells)) and identified differentially expressed genes using the Limma R package⁵¹. The following gene sets were retrieved from RNA-seq datasets: mouse genes related to acetate and butyrate were downloaded from ref.²³ (GEO accession identifier GSE79600) and ref.²⁶ (GEO accession identifier GSE132285), gene sets related to RUNX3 knockout and to HOBIT/BLIMP1 double-knockout from GEO (accession identifiers GSE81885 and GSE70813, respectively). The TOX-related gene set (differentially expressed genes in *Tox*^{-/-} (knockout) versus wild-type P14 cells after 8 days of lymphocytic choriomeningitis virus clone 13 infection) was obtained from ref.¹³. Raw datasets were processed using GREIN DB⁵² and differentially expressed genes were determined using the DESeq2 R package⁴¹. The calcium-signalling-related gene set was downloaded from Mouse Genome Informatics database (<http://www.informatics.jax.org/go/term/GO:0019722>). The human liver tissue-resident memory cell gene signatures were procured from ref.³³ and the IL-15-induced gene set was retrieved from ref.³⁴; these were used for GSEA with single T cell datasets from livers of patients with NASH. GSEA of log₂-transformed fold changes from differentially expressed genes (CXCR6⁺ mice fed a CD-HFD versus CXCR6⁺ mice fed a normal diet, and CXCR6⁺ versus CXCR6⁻ mice fed a CD-HFD) and from single-cell RNA-seq data from CD3⁺ populations of livers from NASH patients were performed using GSEA v.3.0 (ref.⁵³). Fold changes along gene sets were given as an input to the PreRanked tool from GSEA v3.0, which assessed the normalized enrichment score. The normalized enrichment score was considered significant based on an FDR ($q \leq 0.25$).

Statistics

All values are shown as mean $\pm$ s.e.m. Statistical analyses were performed with GraphPad Prism software v.9 using unpaired two-tailed Student's *t*-test, two-way ANOVA and one-way ANOVA analysis with Tukey's, Sidak's or Dunnett's post hoc test for multiple comparisons. $P < 0.05$ was considered significant. No statistical methods were used to predetermine sample size.

Reporting summary

Further information on research design is available in the Nature Research Reporting Summary linked to this paper.

Data availability

All raw gene RNA-seq data from mice have been deposited in the GEO under accession number GSE145104. Single-cell RNA-seq data of fat tissue from patients with NASH can be found in the under accession number EGAS00001004738, and can be accessed at <https://ega-archive.org/access/data-access> upon request to the corresponding author. Single-cell RNA-seq data from livers of patients with steatosis or NASH can be found under accession number GSE159977. All datasets are available from the corresponding author upon request. Source data are provided with this paper.

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Acknowledgements We thank S. Yousuf and M. Qui for technical support in preparing samples for single-cell RNA-seq analysis; P. Sinn for technical support; the personnel (particularly J. Maurer) of the Core Facility Flow Cytometry at Biomedical Center Munich (LMU), for providing the imaging cytometry instrument and analysis software; and the personnel of the Core Facility Cell Sorting of the German Cancer Research Center Heidelberg. This work was supported by the DFG, project number 424926990, to J.P.B., CIBSS EXC-2189 project identifier 390939984 to E.J.P., SFB TRR179 and DZIF Munich site to P.A.K.

Author contributions M.D. and P.A.K. designed the study. M.D., D.P., P.F., A.S., M.L., P.M., F.B., D.I., J.W., S.H., M.A., A.G., A.B., D. Heide, F.M. and S. Reider performed experiments. S.D., R. and A.M.K. undertook bioinformatics analysis. D. Hartmann, N.H., M.S., C.G.-C., T.G., G.S., J.-P.M., M.E., S. Roth, A.T.B., B.M.-S., F.K.-N., R.K., T.B., P.R., V.L., J.-F.D. and M.H. provided tissue samples, essential reagents or mice. R.R., E.J.P., H.T., F.T., J.-F.D., D. Haller, P.J.M., R.H., R.T., D.Z., J.P.B., M.H. and P.A.K. contributed scientific input. M.D. and P.A.K. wrote the manuscript, and all authors contributed to feedback and proofreading.

Competing interests The authors declare no competing interests.

Additional information

Supplementary information The online version contains supplementary material available at <https://doi.org/10.1038/s41586-021-03233-8>.

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Peer review information Nature thanks Paul Klenerman, Ignacio Melero, Daniel Peeper and the other, anonymous, reviewer(s) for their contribution to the peer review of this work. Peer reviewer reports are available.

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Extended Data Fig. 1 | See next page for caption.

Extended Data Fig. 1 | Immune-cell characteristics in livers of mice fed a CD-HFD. a–e, Liver damage after 12 months of feeding a CD-HFD (which induces features of NASH) in *Ja18^{-/-}* and *Cd1d^{-/-}* mice, determined by SALT and NAS with representative images of H&E staining (mice fed a normal diet, *n* = 8; CD-HFD-fed, wild-type mice, *n* ≥ 7; CD-HFD-fed, *Ja18^{-/-}* mice, *n* ≥ 7; CD-HFD-fed, *Cd1d^{-/-}* mice: *n* ≥ 8). Scale bar, 100 μm. Two independent experiments. **f**, Body weight of mice over the course of six months. **g, h**, Numbers of hepatic lymphocytes (**g**) and hepatic CXCR6⁺ T cell populations (**h**) in mice fed a normal diet (*n* = 11) or CD-HFD (*n* = 13). Two independent experiments. **i–k**, Numbers and phenotype of hepatic CD44⁺ CXCR6⁺ and CXCR6⁻ CD8 T cells in NASH mice (*n* ≥ 5). Two independent experiments. **l**, Frequencies of CXCR6⁺GzmB⁺ CD8 T cells in different organs in mice fed a normal diet (*n* = 4)

and NASH mice (*n* = 4). **m–o**, Localization and quantification of hepatic CXCR6⁺PD1^{high} CD8 T cells in mice fed a normal diet (*n* = 3) and NASH mice (*n* = 3), by confocal microscopy with representative images. Scale bar, 200 μm. CD3 (yellow), CD8 (purple), CXCR6 (cyan), PD1 (red) and collagen IV (white). Rectangles highlighted in **m** indicate representative zoomed areas shown in Fig. 1g. ns, not significant; nd, not determined. Exact *P*(**a–d**, **f–l**, **n**, **o**) and *n*(**a–d**, **f**, **i–k**) values are presented in Source Data. **P* < 0.05, ***P* < 0.01, ****P* < 0.001, *****P* < 0.0001. Two-way ANOVA with Sidak's (**g**, **h**, **l**) or with Tukey's multiple comparison test (**j**, **k**), one-way ANOVA with Tukey's (**a–d**) or with Dunnett's multiple comparison test (**f**) and unpaired two-tailed *t*-test (**i**, **n**, **o**). In **a–d**, **f–l**, **n**, **o**, data are mean ± s.e.m., error is reported as s.d.

fed a normal diet or CD-HFD. Coefficient of determination (R^2) and statistical significance (P value) were determined using Pearson's correlation. **e, f**, CXCR6 and PD1 expression levels in hepatic CD8 T cells in mice fed CD-HFD (**e**) or a high *trans*-fat Western diet (WD) (**f**) ($n = 5$) for 12 months compared to mice fed a normal diet ($n = 5$). **h, i**, Liver damage (SALT and NAS) in mice from **f, j, k**, CXCR6 and PD1 expression levels in hepatic CD8 T cells in *ob/-* and *ob/ob* mice after 4 months of chow feeding or mice fed a normal diet (*ob/-*, $n = 6$; *ob/ob*, $n = 4$). **l, m**, Liver damage (SALT and NAS) in mice from **j**. Exact P values (**b, f-m**) are presented in Source Data. * $P < 0.05$, ** $P < 0.01$, **** $P < 0.0001$ (unpaired two-tailed t -test (**f-m**). In **b, f-m**, data are mean $\pm$ s.e.m., errors are shown as s.d.

Extended Data Fig. 3 | Transcriptome analysis of hepatic CXCR6⁺ CD8 T cells from NASH mice. **a**, Volcano plot showing differentially expressed genes between CXCR6⁺ CD8 T cells from NASH mice (fed a CD-HFD) compared to healthy mice (fed a normal diet). **b**, GSEA for distinct pathways from **a**. **c**, Flow cytometry analysis of protein levels of transcription factors in hepatic CXCR6⁺ and CXCR6^{neg} CD8 T cells from healthy and NASH mice ($n \geq 5$). **d**, Binding analysis for regulation of transcription (BART) to predict transcription-factor activity in hepatic CXCR6⁺ compared to CXCR6^{neg} CD8 T cells in NASH mice. **e**, Transcription-factor network analysis created from differentially expressed genes (**a**) and BART. **f**, GSEA for transcription-factor-dependent genes from **a**. **g-i**, Flow cytometry analysis of FOXO1 expression in hepatic CXCR6⁺ and

CXCR6^{neg} CD8 T cells (**g, h**) or CD4 T cells (**i**) from mice fed a normal diet and NASH mice ($n \geq 6$). **j, k**, Image-stream analysis of expression and localization of FOXO1 in different T cell populations from spleen and liver of wild-type mice with representative images. Tn, naive T cells; Tcm, central memory T cells; Tem, effector memory T cells. **l**, Similarity score to determine nuclear localization of FOXO1 and surface localization of CD44 ($n = 5$). **m, n**, FOXO1 expression in hepatic CD44⁺ CD8 T cells in mice fed a Western diet ($n = 5$) and *ob/ob* or *ob/ob* mice ($n = 4$). PI, propidium iodide. Exact *P* and *n* values (**c, g-i**) are presented in Source Data. **P* < 0.05, ***P* < 0.01, *****P* < 0.0001. One-way ANOVA (**k, l**), two-way ANOVA with Tukey's multiple comparison test (**c, i**) and unpaired two-tailed *t*-test (**h, m, n**). In **c, h, i, k-n**, data are mean $\pm$ s.e.m., errors are shown as s.d.

Extended Data Fig. 4 | Hepatic CXCR6⁺FOXO1^{low} CD8 T cell frequency correlates with liver damage in NASH mice and patients with NASH.

a–c, Hepatic CXCR6⁺FOXO1^{low} CD8 T cells at 6, 9 and 12 months of feeding a CD-HFD, and correlation of FOXO1 expression levels in hepatic CD44⁺CD8⁺ T cells with frequency of hepatic CXCR6⁺PD-1^{high} of CD44⁺CD8⁺ T cells from mice fed a CD-HFD, and with NAS. Four independent experiments. **d, e**, CXCR6 and CX₃CR1 expression levels in sorted hepatic CD44⁺CXCR6⁺CD8⁺ T cells from healthy mice fed a normal diet ($n = 5$) after 18 h of treatment with the FOXO1 inhibitor AS1842856. **f**, CXCR6 expression levels after FOXO1 overexpression (transduction with pMP71-Foxo1-eGFP or pMP71-eGFP) in sorted CXCR6⁺CD8⁺ T cells ($n = 3$). **g, h**, Correlation of sALT and NAS with FOXO1 expression in hepatic CD44⁺CD8⁺ T cells from **b**. **i**, Correlation of FOXO1 expression in liver CD69⁺CD8⁺ T cells with sALT in patients with NASH ($n = 15$ in total: $n = 12$ NASH;

$n = 3$ normal liver tissue (NAS = 0)). **j**, CXCR6 expression quantified by qPCR with reverse transcription (RT-qPCR) in liver tissue of patients with NASH (NAS ≥ 5) (unaffected tissue, $n = 6$; tissue from patients with NASH, $n = 7$). **k, l**, Correlation of relative CXCR6 expression with sALT and NAS in patients with NASH (unaffected individuals, $n = 6$; patients with NASH, $n = 10$). **m**, Correlation of sALT with numbers of cells that are positive for CXCR6 per mm² of liver tissue in patients with NASH ($n = 18$). Scale bar, 100 μ m. Representative image of in situ hybridization for CXCR6 mRNA in liver of patients with low or high sALT. Exact P (**e, f, j**) and n (**b, c, g, h**) values are presented in Source Data. * $P < 0.05$, ** $P < 0.01$, **** $P < 0.0001$. Coefficient of determination (R^2) and statistical significance (P value) were determined using Pearson's correlation (**b, c, g–i, k–m**). Unpaired two-tailed (**f, j**) and paired two-tailed t -test (**e**). In **e, f, j**, data are mean $\pm$ s.e.m., error is reported as s.d.

Extended Data Fig. 5 | Increased levels of IL-15 in serum and liver tissue of NASH mice correlate with liver damage and numbers of hepatic CXCR6⁺ CD8 T cells. **a**, Heat map of fold changes for cytokine and chemokine serum concentrations in NASH mice (fed a CD-HFD) (*n* = 8) compared to healthy mice (fed a normal diet) (*n* = 6) after 12 months of feeding. Values above 2.5 are red. Asterisk denotes values that are significantly different. Exact *P* and *n* values are presented in Source Data. **b**, Cytokine and chemokine expression quantified by RT-qPCR in the liver tissue of mice fed a normal diet (*n* = 5) and NASH mice (*n* = 9). **c**, Flow cytometry analysis of the increase in CXCR6 expression after incubation of sorted hepatic CD44⁺ CXCR6⁺ CD8 T cells with cytokines for 24 h (*n* = 13). **d**, Flow cytometry analysis of FOXO1 expression 24 h after IL-15 stimulation of hepatic CD44⁺ CD8 T cells (*n* = 10). **e–g**, Flow cytometry analysis of FOXO1 expression in human CD8 T cells at 5 d after IL-15 stimulation (15 ng ml⁻¹) and frequencies of CXCR6⁺ CD69⁺ cells (*n* = 5). **h**, PD1, GzmB and CXCR3 expression by mouse hepatic CXCR6⁺ CD8 T cells 24 h after IL-15 stimulation (*n* = 9). **i**, Percentage of CXCR6-expressing CD8 T cells in

sorted, splenic CD44⁺ CD8 T cells 2 d after stimulation with IL-15 (10 ng ml⁻¹) and/or IL-21 (10 ng ml⁻¹) (*n* = 5). Two independent experiments. **j**, **k**, IL-15 serum concentrations in *ob/ob* (*n* = 4) and hyperphagic, leptin-deficient *ob/ob* mice (*n* = 4) fed a chow diet for 4 months, in mice fed a normal diet (*n* = 4) or a high trans-fat Western diet for 12 months (*n* = 5). **l–n**, Correlation of IL-15 serum levels with sALT, NAS and FOXO1 expression in hepatic CD44⁺ CD8 T cells of NASH mice (6 months, *n* = 4; 9 months, *n* = 7; 12 months, *n* = 13). Four independent experiments. **o–q**, Correlation of relative hepatic IL-15 gene expression levels with sALT, NAS and CXCR6 gene expression in patients with NASH and in individuals with normal liver (normal, *n* = 6; NASH, *n* = 10). Exact *P* (**a–d**, **f–k**) and *n* (**a**) values are presented in Source Data. **P* < 0.05, ***P* < 0.01, ****P* < 0.001, *****P* < 0.0001. Coefficient of determination (*R*²) and statistical significance (*P* value) was determined using Pearson's correlation (**l–q**). One-way ANOVA with Tukey's (**i**) or Dunnett's multiple comparison test (**c**), and unpaired two-tailed *t*-test (**a**, **b**, **d**, **f–h**, **j**, **k**). In **a–d**, **f–k**, data are mean ± s.e.m., error is reported as s.d.

Extended Data Fig. 6 | Single-cell RNA-seq of CD3⁺ T cells from livers of patients with steatosis or NASH. **a**, UMAP plot visualization of hepatic CD3⁺ T cell clusters detected in livers from patients with steatosis ($n=4$) or NASH ($NAS \geq 5$) ($n=3$). **b**, UMAP plot visualization of hepatic CD3⁺ T cell clusters and of selected genes detected in patients with NASH ($NAS \geq 5$) ($n=3$). **c**, Frequencies of CXCR6⁺ and PDCD1⁺ cells in CD8 T cells in the *RGS1*⁺, *PLCG2*⁺ and *ITGAI1*⁺ cluster in livers from patients with steatosis or NASH. **d**, Differentially expressed genes in the *CD8RGS1*⁺ cluster between livers of patients with steatosis or NASH. **e**, Flow cytometry analysis of FOXO1 expression in CD69⁺ CD8 T cells from livers of unaffected individuals ($n=3$) and livers of patients with NASH ($n=5$).

$P=0.0088$, two-tailed t -test, data are mean $\pm$ s.e.m. **f**, UMAP plot visualization of hepatic CD3⁺ T cell clusters detected in patients with NASH from fat tissue ($n=2$). **g**, Jaccard similarity score between tissue-resident CD8 T cell populations in fat and liver of patients with NASH ($n=2$). **h**, Correlation of mouse RNA-seq dataset from CXCR6⁺ compared to CXCR6⁺ CD8 T cells in mice fed a CD-HFD with human single-cell RNA-seq dataset from the CD8 *GNLY*⁺ *FGFBP2*⁺ cluster. The solid line is the linear regression, and the grey area shows the 95% confidence intervals. Coefficient of determination (R^2) and statistical significance (P value) were determined using Spearman's correlation.

Extended Data Fig. 7 | Anti-CD122 application in NASH mice leads to depletion of CXCR6⁺Gzmb⁺ CD8 T cells and amelioration of liver damage. **a**, CD122 expression on splenic CD44⁺CD8 T cells after stimulation with IL-15 (10 ng ml⁻¹) for 24 h ($n=5$). **b**, CD122 expression on hepatic CXCR6⁺CD8 T cells ($n=8$). Two independent experiments. **c–e**, Frequencies and cell numbers of CXCR6⁺PD1⁺ and CXCR6⁺Gzmb⁺CD8 T cells after anti-CD122-treatment for 2 weeks in NASH mice (fed CD-HFD for 12 months) (CD-HFD + immunoglobulin (Ig), $n=6$; CD-HFD + anti-CD122, $n=6$). **f**, sALT in NASH mice from **e** before and after anti-CD122 treatment for 2 weeks. Unpaired two-tailed t -test of sALT

values between groups before treatment, $P=0.7137$. **g–k**, Numbers of CX₃CR₁CD8 T cells, CD44⁺CD4 T cells, natural killer T cells and natural killer (NK) cells in the liver after anti-CD122 antibody application from mice in **e**. **l, m**, No change in phenotype of natural killer cells in NASH mice. Dot plots are representative of ≥ 5 mice. Exact P values (**a, b, e–m**) are presented in Source Data. * $P<0.05$, ** $P<0.01$, *** $P<0.001$, **** $P<0.0001$. One-way ANOVA with Tukey's multiple comparison test (**e, g–k**) and paired (**a, f**) and unpaired (**b, l, m**) two-tailed t -test. In **a, b, e–m**, data are mean $\pm$ s.e.m., error is reported as s.d.

Extended Data Fig. 8 | See next page for caption.

Extended Data Fig. 8 | IL-15-treated CXCR6⁺ CD8 T cells become auto-aggressive after exposure to acetate. **a**, Auto-aggressive activity of IL-15-treated CXCR6⁺ FOXO1^{low} CD8 T cells against hepatocytes ($n = 3$). **b**, GzmB-expression levels in CXCR6⁺ CD8 T cells exposed for 24 h to different concentrations of acetate in vitro ($n = 4$). **c–e**, Fluorospot assay with representative images to quantify TNF and IFN γ release from IL-15-stimulated CXCR6⁺ and CXCR6⁺ CD8 T cells after acetate exposure and from peptide-specific stimulation of OT1 CD8 T cells ($n = 5$). **f**, GSEA for differentially expressed genes from hepatic CXCR6⁺ and CXCR6⁺ CD8 T cells of mice fed a CD-HFD compared to in vitro-generated memory CD8 T cells exposed to acetate or butyrate. **g**, Acetate concentrations in supernatants of primary mouse hepatocytes after exposure to palmitate for 24 h (250 μ M) or with an agonistic trimeric FasL (25 ng ml⁻¹) to induce cell death ($n = 6$). **h**, Cytotoxic activity of acetate-exposed polyclonal CD8 T cells after FOXO1 inhibition (AS1842856, 100 nM) against hepatocytes ($n = 3$) and quantification. **i**, GzmB expression in acetate-exposed, GFP-expressing CD8 T cells after retroviral transduction with pMP71-GFP-Foxo1 or pMP71-GFP ($n = 3$). **j**, **k**, sALT at day 2 after adoptive transfer of IL-15-stimulated or FOXO1-inhibited and acetate-exposed CD8 T cells ($n \geq 3$). **l**, Calculation of liver-damage-inducing potential per cell for auto-aggressive CD8 T cells ($n = 8$). **m**, Surface LAMP1 expression on CXCR6⁺ CD8 T cells after a 24-h exposure to acetate (15 mM) ($n = 9$). **n**, **o**, Liver damage quantified by NAS and sALT in *Prf1*^{-/-} and littermate control mice after CD-HFD

feeding for 1 year. **p**, ICAM1 expression on hepatocytes in coculture with IL-15-stimulated, acetate-exposed CXCR6⁺ CD8 T cells ($n = 7$). **q**, Liver immunohistochemistry with representative images and quantification of ICAM1 clusters per mm² of liver tissue in mice fed a normal diet ($n = 5$), NASH mice ($n = 6$) and NASH mice treated with anti-CD122 antibody ($n = 6$). Scale bar, 100 μ m. **r**, Representative microscopic images of IL-15-stimulated, acetate-exposed CXCR6⁺ CD8 T cells in contact with hepatocytes and quantification of the distance. Arrows indicate polarized LFA1 orientation on T cell interacting with hepatocyte. Scale bars, 30 μ m (left), 15 μ m (right). **s**, Auto-aggressive activity of IL-15-treated CXCR6⁺ CD8 T cells against hepatocytes in presence of acetate (15 mM) or TNF (5 ng ml⁻¹) and anti-LFA1 antibodies ($n = 3$). **t**, sALT at day 2 after adoptive transfer of auto-aggressive CD8 T cells and injection of anti-LFA1 (100 μ g per mouse) or control immunoglobulin. Two independent experiments. **u**, **v**, *TNF* expression in livers from unaffected individuals ($n = 6$) or patients with NASH ($n = 7$) (**u**) and correlation with sALT (healthy, $n = 6$; NASH, $n = 10$) (**v**). Cytotoxic activity results are representative of at least three independent experiments. Exact P (**a**, **b**, **d**, **e**, **g–v**) and n (**n**, **o**, **t**) values are presented in Source Data. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$, **** $P < 0.0001$. Two-way ANOVA with Sidak's multiple comparison test (**d**, **e**), one-way ANOVA with Tukey's (**j**, **k**, **s**, **t**) or with Dunnett's (**b**, **g**, **h**) multiple comparison test and unpaired two-tailed t -test (**a**, **i**, **l–n**, **p**, **r**, **u**). In **a**, **b**, **d**, **e**, **g–v**, data are mean $\pm$ s.e.m., error is reported as s.d.

Extended Data Fig. 9 | Metabolic activation of CXCR6⁺ CD8 T cells triggers auto-aggression. **a**, Phosphorylated proteins downstream of TCR signalling in IL-15-stimulated, acetate-exposed CXCR6⁺ CD8 T cells. **b**, Glycolysis in auto-aggressive CXCR6⁺ CD8 T cells (n = 3). **c**, Calcium influx in IL-15-stimulated CXCR6⁺ and CXCR6⁻ CD8 T cells exposed to metabolites, cytokines or TLR ligands (n = 4). **d**, **e**, Extracellular ATP in primary mouse hepatocyte supernatant after 24 h treatment with palmitate (250 μ M) or agonistic trimeric FasL, or coculture with auto-aggressive CD8 T cells (n = 3). **f**, **g**, Metabolite detection in liver tissue of mice fed a normal diet or CD-HFD by matrix-assisted laser desorption and ionization imaging. Scale bar, 500 μ m (**f**). Exact minima, maxima, centres, bounds of box and whiskers and percentiles are presented in Source Data. **h**, P2RX7 expression in IL-15-stimulated CD44⁺ CXCR6⁺ CD8

T cells. **i**, **j**, Calcium influx in CXCR6⁺ CD8 T cells after incubation with NAD (1 μ M) (n = 5) or supernatants from **d** (n = 3) for 30 min. **k**, Representative cytotoxicity result of CD8 T cell auto-aggression in the presence of P2RX7 or PANX1 inhibition (n = 3). **l**, sALT at day 2 after transfer of auto-aggressive CD8 T cells and anti-P2RX7 nanobody (50 μ g per mouse). Two independent experiments. **m**, Representative cytotoxicity result of antigen-specific, IL-15-stimulated and acetate-exposed CXCR6⁺ OT1 CD8 T cells against S8L-loaded hepatocytes. Exact P (**a**–**e**, **i**, **j**, **l**) and n (**a**, **l**) values are presented in Source Data. *P < 0.05, **P < 0.01, ****P < 0.0001. Two-way ANOVA with Tukey's multiple comparison test (**c**, **i**, **j**), one-way ANOVA with Tukey's (**b**, **e**, **l**) or with Dunnett's (**d**) multiple comparison test and unpaired two-tailed t-test (**a**). In **a**–**e**, **i**, **j**, **l** data are mean $\pm$ s.e.m., error is reported as s.d.

Extended Data Fig. 10 | Auto-aggressive human CD8 T cells. **a**, Auto-aggression of allogeneic and autologous CD8 T cells against primary human hepatocytes after incubation with IL-15 (effector-to-target ratio of 3:1 ($n=3$)). **b**, Auto-aggression by IL-15-treated human CD8 T cells exposed to acetate ($n=3$). **c**, Acetate levels in livers from unaffected individuals or patients with NASH ($n=3$). **d**, Auto-aggression of human CD8 T cells in presence of anti-TNF ($10 \mu\text{g ml}^{-1}$) or anti-LFA1 ($10 \mu\text{g ml}^{-1}$) ($n=3$). **e**, **f**, Auto-aggression against MHC-I-deficient K562 cells by human CD8 T cells exposed to IL-15 (**e**), and at different effector-to-target ratios (**f**) ($n=6$). **g**, Calcium influx in IL-15-treated human CXCR6⁺CD69⁺ CD8 T cells measured with a calcium sensor dye before or after coculture with K562 cells and quantification ($n=6$). **h**, **i**, Auto-aggressive killing of K562 cells by IL-15-treated CD8 T cells in presence of anti-FasL ($10 \mu\text{g ml}^{-1}$), anti-NKG2D ($10 \mu\text{g ml}^{-1}$) or EGTA (4 mM) at effector-to-target ratio of 10:1 ($n=6$). **j**, Mean numbers of CD8 T cells adjacent

to cleaved CASP3⁺ hepatocytes in mice fed a normal diet and NASH mice ($n=3$). **k**, **l**, Surface FasL and TRAIL expression of mouse CXCR6⁺ CD8 T cells after 1 h exposure to ATP (200 μM). **m**, Caspase activity in primary mouse hepatocytes cocultured with auto-aggressive CD8 T cells and anti-FasL ($10 \mu\text{g ml}^{-1}$) ($n=3$). **n**, Mouse CD8 T cell auto-aggression in presence of inhibitors against apoptosis (ZVAD-FMK, $10 \mu\text{M}$) or necroptosis (Nec-1s, $30 \mu\text{M}$) ($n=4$). **o**, sALT at day 2 after adoptive transfer of IL-15-treated, acetate-exposed CD8 T cells and anti-FasL injection (100 μg per mouse). Two independent experiments. Exact P (**a–o**) and n (**k**, **l**, **o**) values are presented in Source Data. * $P<0.05$, ** $P<0.01$, *** $P<0.001$, **** $P<0.0001$. Two-way ANOVA with Sidak's (**a**) and Tukey's (**f**, **g**) multiple comparison test, one-way ANOVA with Tukey's multiple comparison test (**b**, **d**, **e**, **h**, **i**, **m–o**) and unpaired two-tailed t -test (**c**, **j–l**). In **a–o**, data are mean $\pm$ s.e.m., error is reported as s.d.

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Software and code

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Data collection

Data for RNA-sequencing analysis was generated from hepatic CD8 T cells of mice fed a normal diet or choline-deficient high fat diet for 12 months.

Data for single-cell RNA-sequencing analysis was generated from hepatic CD45+ cells of steatotic and NASH patients undergoing bariatric surgery.

Data for single-cell RNA-sequencing analysis was generated from CD45+ CD3+ cells of from fat tissue of NASH patients undergoing bariatric surgery.

Genesets from public available sources used for GSEA are described in the method section.

Following machines for data collection were used: SP6800, SA3800 and SH800 (Sony Biotechnology); Seahorse XF 96 Analyser (Agilent Technologies); Luminex 200 Instrument System (Thermo) ; xCelligence RTCA MP device (ACEA Biosciences); Leica Biosystems Bond MAX; Tecan Infinite M1000 (Tecan); Leica confocal microscope TCS SP8; TimsTOF Flex mass spectrometer (Bruker); Immunospots® S6 ULTIMATE Analyzer; LightCycler 480 (Roche); ViiA7 (Applied Biosystems); NextSeq 500 (Illumina); HiSeq 3000 and NovaSeq6000 (Illumina); Amnis ImageStream (Millipore); Reflotron®plus system (Roche); Fuji DRI-CHEM NX500i (Fujifilm)

Data analysis

Following machines for data analysis were used: Drop-seq 1.12, DESeq2 v2.1.28.1 R package; Seurat v3.2.2 R package; BART v1.1, igraph v1.2.6 R package; Cytofit v1.11.3 R package; pheatmap v1.0.12 R package; ggpubr v0.4.0 R package; ggplot2 v3.3.2 R package; Graph Pad Prism v9 (Graph-Pad Software); FlowJo v10.8; Arivis Vision 4D (Arivis); Limma v3.44.3 R package; Cytoscape v3.7.1; MSigDB v6.2; GSEA v4.0.3; Sony SP Spectral Analyzer Software 2.0.2.14140 (Sony Biotechnology); Sony SA Software 2.0.2.04040 (Sony Biotechnology); SH Software V1.8.2 (Sony Biotechnology); Imaris software (Version 9.5, Bitplane); SciLS Lab (Bruker Daltonics); Seahorse XF 96 Analyzer Software v1.4.2 (Agilent Technologies); CTL Immunospot 5.4 Professional DC Software (CTL Europe); IDEAS software v6.2; RTCA Software Pro; Luminex 200

instrument system software; for further details please see the materials and methods section.

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Data from murine RNAseq are deposited at GEO accession number: GSE145104.

Single-cell RNA-sequencing data of CD45+ cells from liver tissue of steatotic and NASH patients: GSE159977.

Single cell RNA-sequencing data of fat tissue from NASH patients: EGAS00001004738.

All other datasets generated during the current study are available from the corresponding authors on reasonable request.

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Sample size	Pilot experiments and previous published results (Wolf et al, 2014; Malehmir et al., 2019) were used to estimate the sample size such that appropriate statistical tests could yield significant results. The exact n numbers used in the study are indicated in the respective figure legends.
Data exclusions	Every mouse fed a normal diet or a choline-deficient high fat diet was included in the analysis. The complete dataset on any individual human or murine sample for flow cytometry was excluded if any obvious problem during sample processing occurred, i.e. more than 70 percent dead cells identified by live dead staining.
Replication	All experiments were repeated and validated as stated in the respective figure legends.
Randomization	Murine experiments: 6-week-old C57Bl/6 mice were randomly allocated into different groups and were fed with appropriated diet or treatment regimens. Human experiments: As described in the figure legends, statistical analysis was corrected for the covariate age.
Blinding	Murine experiments were rated in a blinded manner. Further analysis was performed unblinded because the type of diet (ND and CD-HFD) could be identified by the researcher on the basis of the colour of the food (ND=brown, CD-HFD=green). Immunohistochemistry, confocal microscopy and serum parameter analysis was performed in a blinded fashion with code-labelling. Every experiment was performed using several control samples and the applied analysis strategy (especially FACS gating strategy) was identical for every single sample. Human experiments: During analysis, patient data underwent pseudonymisation and was blinded to the analyser.

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Antibodies

Antibodies used

Following mouse antibodies for flow cytometry were purchased from BioLegend: anti-CD4 (PM4-5), anti-CD8 (53-6.7), anti-CD44 (IM7), anti-CD69 (H1.2F3), anti-CD178 (FasL, MFL3), anti-CD183 (CXCR3-173), anti-CD186 (CXCR6, SA051D1), anti-CD253 (TRAIL, N2B2), anti-CD279 (PD-1, 29F.1A12), anti-CX3CR1 (SA011F11), iFN γ (XM61.2), anti-KLRG1 (2F1), anti-NK1.1 (PK136), anti-TCR β (H57-597). Following mouse antibodies for flow cytometry were purchased from ThermoFisher: anti-Blimp1 (5E7), anti-Bcl-2 (100), anti-CD3 (17A2), anti-CD11a (1217), anti-CD45.1 (A20), anti-CD49a (HMa1), anti-CD54 (ICAM-1, YN1/1.7.4), anti-CD62L (MEL-14), anti-CD107a (LAMP1, eBioD4B), anti-CD122 (IL2rb, 5H4), anti-EOMES (Dan11mag), anti-Gzmb (GB11), anti-P2XR7 (1F11), anti-pAkt (SDRNR), anti-pZAP70 (h3kobu5), anti-plck (SRRCHA), anti-T-bet (eBio4B10), anti-TNF (MP6-XT22), TOX (TXRX10). Anti-Foxo1 (C29H4) was purchased from Cell Signaling.

Following human antibodies for flow cytometry were used: anti-CD3 (SK7, BioLegend), anti-CD8 (HIT8a, BioLegend), anti-CD45 (HI-30, BioLegend), anti-CD69 (FN50, BioLegend), anti-CD186 (K041E5, BioLegend), anti-Gzmb (GB11, ThermoFisher), anti-Foxo1 (C29H4, Cell Signaling).

Detection of cytokine concentrations in serum was performed with a 23-plex bead array (Bio-Rad, #M60009RDPD) and with a Bio-Plex against CXCL16 (Bio-Rad, #12002257) or with enzyme-linked immunosorbent assay (ELISA) for IL-15 (Peprotech, #900-K188) or IL-21 (Peprotech, #900-K368).

Following murine antibodies were used for confocal microscopy of CXCR6 and PD-1: anti-CD3 (Sony, clone 17A2, #1101045, 1 μ g/ml), anti-CD8 (eBioscience, clone 53-6.7, #56-0081-82, 2 μ g/ml), anti-CXCR6 (Sony, clone SA051D1, #1355545, 1 μ g/ml), anti-PD-1 (R&D Systems, polyclonal, #AF1021, 1 μ g/ml) and anti-Collagen IV (Abcam, polyclonal, #Ab6586, 0.5 μ g/ml). Unconjugated primary antibodies were stained with the following secondary antibodies over night: donkey anti-rabbit IgG (BioLegend, clone Poly4064, #406404, 1 μ g/ml) and donkey anti-goat IgG (Jackson Immunology, polyclonal, #705-587-003, 1 μ g/ml).

rat anti-mouse CD8 (1:200, 45M15, ThermoFisher), rabbit anti-mouse cleaved caspase-3 (1:300, D3E9, Cell Signaling Technology), goat anti-mouse collagen IV-a1 (#nb120-6586, 1:30, polyclonal, Novus Biologicals). The following secondary Abs were used for staining: Alexa Fluor 488 chicken anti-rat IgG (H+L) (#A-21470, ThermoFisher, 1 μ g/ml), Alexa Fluor 568 goat anti-rabbit IgG (H+L) (#A-11011, ThermoFisher, 2 μ g/ml), Alexa Fluor 647 donkey anti-goat IgG (H+L) (#A-21447, ThermoFisher, 2 μ g/ml).

Following murine antibodies were used for multiplex immunohistochemistry: anti-PD-1 (1/200, R&D Systems, #AF1021), anti-CD8a (1/200, ThermoFisher #14-0808-82), anti- α SMA (1/200, Agilent #M085129-2), anti-IBA1 (1/1,000, EMD Millipore #MABN92), anti-CK19 (1/200, DSHB #TROMA-III). TROMA-III was deposited to the DSHB by Kemler, R. (DSHB Hybridoma Product TROMA-III).

Following murine antibodies were used for immunohistochemistry: anti-ICAM-1, 1:1500, clone 3E2B, #MA5405, ThermoFisher

Following murine and human antibodies were used for CD8 T cell cytotoxicity assays: TNF (ThermoFisher, #PMC3013), IL-6 (Miltenyi Biotec, #130-094-065), IL-1 (Peprotech, #211-11B), anti-LFA1 (mouse: BioXcell, M17/4, BE0006, 10 μ g/ml; human: BioXcell, #BE0192, R7-1, 10 μ g/ml), anti-mouse MHC1 (BioXcell, M1/42.3.9.8, #BE0077, 10 μ g/ml), anti-mouse PD-1 (BioXcell, RMP1-14, #BE0146, 10 μ g/ml), anti-mouse TNFR1 (BioLegend, 55R-593, #113102, 10 μ g/ml), anti-mouse TNFR2 (BioLegend, TR75-32.4, #113202, 10 μ g/ml), anti-human TNF (BioLegend, MAb11, #502901), anti-mouse iFN γ (BioLegend, XM61.2, #505801, 10 μ g/ml), anti-FasL (mouse: BioXcell, MFL3, #BE0319, 10 μ g/ml; human: BioLegend, NOK-1, #306415, 10 μ g/ml), anti-human NKG2D (BioLegend, 1D11, #320813, 10 μ g/ml)

Validation

All antibodies described in the previous section underwent quality control by the companies.

BioLegend tested the quality of their antibodies for flow cytometry by using the following protocols: intracellular (True-Nuclear™ Transcription Factor Staining Protocol, <https://www.biolegend.com/protocols/true-nuclear-transcription-factor-staining-protocol-for-5ml-tubes/4267/>) and surface (<https://www.biolegend.com/protocols/cell-surface-flow-cytometry-staining-protocol/4283>). Neutralization capacity of anti-human TNF (Abiamunits V, et al. 2010. Eur. J. Immunol. 40:2891), anti-human FasL (Kayagaki N, et al. 1995. J. Exp. Med. 182:1777) and anti-human NKG2D (Groh V, et al. 2001. Nature Immunol. 2:255) was reported and not tested by BioLegend in house.

ThermoFisher: all antibodies for surface and intracellular targets have been tested by flow cytometric analysis of mouse splenocytes or human peripheral blood mononuclear cells (e.g. mouse CD3: <https://www.thermofisher.com/antibody/product/CD3-Antibody-clone-17A2-Monoclonal/11-0032-82>), e.g. mouse EOMES: <https://www.thermofisher.com/antibody/product/EOMES-Antibody-clone-Dan11mag-Monoclonal/53-4875-82>, e.g. human Gzmb: <https://www.thermofisher.com/antibody/product/Gzmb-Antibody-clone-GB11-Monoclonal/12-8899-41>).

Cell Signaling: Foxo1 antibody was tested for flow cytometry by using Jurkat cells and anti-rabbit (H+L), F(ab')₂ Fragment (PE Conjugate, #8885) as a secondary antibody with the following protocol: <https://www.cellsignal.com/products/primary-antibodies/foxo1-c29h4-rabbit-mab/2880>.

Peprotech: validation of IL-15 and IL-21 for ELISA application is described here: <https://www.peprotech.com/de/anti-murine-il-15> and <https://www.peprotech.com/de/anti-murine-il-21>

R&D: validation of PD-1 for IHC is described here: https://www.rndsystems.com/products/mouse-pd-1-antibody_af1021

BioXcell: validation of binding specificity for each blocking antibody was performed by western blotting of cell lysates of transfected cell lines with cDNAs or by flow cytometry (e.g. anti-PD-1: <https://bxcell.com/product/invivomab-anti-m-pd-1/>, anti-LFA1: C57BL/6 mouse splenic secondary cytotoxic T cells)

Abcam: description of the validation of anti-collagen IV antibody can be found here: <https://www.abcam.com/collagen-iv-antibody-ab6586.html>.

Novus biologicals: validation of antibody specificity of anti-mouse collagen IV was performed by genetic strategies: https://www.novusbio.com/products/collagen-iv-antibody_nb120-6586

Eukaryotic cell lines

Policy information about cell lines

Cell line source(s)

K562 cells were obtained from ATCC <https://www.lgcstandards-atcc.org/products/all/CCL-243.aspx>
Retroviral packaging cell line Plat-Eco were obtained from Cell Biolabs <https://www.cellbiolabs.com/platinum-e-plat-e-retroviral-packaging-cell-line>

Authentication

The cells were authenticated by the vendor and no further authentication was performed in the laboratory

Mycoplasma contamination

All cell lines were routinely tested negative for mycoplasma contamination

Commonly misidentified lines (See [ICLAC](#) register)

No commonly misidentified cell lines were used

Palaeontology and Archaeology

Specimen provenance

Provide provenance information for specimens and describe permits that were obtained for the work (including the name of the issuing authority, the date of issue, and any identifying information).

Specimen deposition

Indicate where the specimens have been deposited to permit free access by other researchers.

Dating methods

If new dates are provided, describe how they were obtained (e.g. collection, storage, sample pretreatment and measurement), where they were obtained (i.e. lab name), the calibration program and the protocol for quality assurance OR state that no new dates are provided.

☐ Tick this box to confirm that the raw and calibrated dates are available in the paper or in Supplementary Information.

Ethics oversight

Identify the organization(s) that approved or provided guidance on the study protocol, OR state that no ethical approval or guidance was required and explain why not.

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Animals and other organisms

Policy information about [studies involving animals](#); [ARRIVE guidelines](#) recommended for reporting animal research

Laboratory animals	<p>- C57Bl/6 mice were purchased from Charles River (Sulzfeld, Germany). H-2Kb-restricted T-cell receptor (TCR) transgenic CD45.1+ OT-1 mice were bred and maintained under specific pathogen-free (SPF) conditions in the centre for preclinical research, School of Medicine, TUM, according to the guidelines of the Federation of Laboratory Animal Science Association. Male mice between the ages of 6 and 10 weeks were used in experiments. All experiments with these mice were approved by Regierung Oberbayern (55.2-1-54-2532-60-14)</p> <p>- 6-week-old male C57Bl/6, Prf1^{-/-}, Cd1d^{-/-}, Ja18^{-/-} and Tcrb^{-/-}, Tcrd^{-/-} mice were fed with a normal diet, a choline-deficient high fat diet or western diet with trans fat ad libitum and housed at the at German Cancer Research Center (DKFZ) according to the guidelines of the Federation of Laboratory Animal Science Association. All experiments with these mice including interventional studies were performed in accordance to German Law (g129/16).</p>
Wild animals	No wild animals were used in this study
Field-collected samples	No field-collected samples were used in this study
Ethics oversight	All mice were kept according to the guidelines of the Federation of Laboratory Animal Science Association and experiments were authorized by permission of the Regierung of Oberbayern and the animal reserach committee at the DKFZ g129/16 and 55.2-1-54-2532-60-14.

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Human research participants

Policy information about [studies involving human research participants](#)

Population characteristics	Patients with NAFLD/NASH were admitted for minilaroscopy-guided liver biopsy or bariatric surgery at the University Hospital Hamburg Eppendorf, at the University Clinic for Visceral Surgery and Medicine (Inselspital), at the Medical University Innsbruck, at the Department of General and Visceral Surgery (University of Freiburg) or at the Heidelberg University Hospital. The patients were recruited into the study after they gave written and informed consent.
Recruitment	Consecutive patients presenting to the outpatient's department at the University hospitals of Heidelberg, Freiburg, München, Hamburg, Bern and Innsbruck were included into the study after they gave written informed consent.
Ethics oversight	All experiments with human liver samples were approved by the ethics committee of Hamburg, Germany (PV5184 and PV5187), Innsbruck, Austria (AN2017-0016 369/4.21) and Bern, Switzerland (2020-00852). Single cell RNAseq-analysis of CD3 + lymphocytes from human liver tissue and mesenterial fat tissue was approved by the Medical Ethics Committee of Heidelberg University (S-629/2013) and the Ethics Committee of Freiburg University (544/17). Experiments with normal human liver samples for isolating primary human hepatocytes were approved by the ethics committee at TUM (232/19S).

Clinical data

Policy information about [clinical studies](#)

All manuscripts should comply with the ICMJE [guidelines for publication of clinical research](#) and a completed [CONSORT checklist](#) must be included with all submissions.

Clinical trial registration	<i>Provide the trial registration number from ClinicalTrials.gov or an equivalent agency.</i>
Study protocol	<i>Note where the full trial protocol can be accessed OR if not available, explain why.</i>
Data collection	<i>Describe the settings and locales of data collection, noting the time periods of recruitment and data collection.</i>
Outcomes	<i>Describe how you pre-defined primary and secondary outcome measures and how you assessed these measures.</i>

Dual use research of concern

Policy information about [dual use research of concern](#)

Hazards

Could the accidental, deliberate or reckless misuse of agents or technologies generated in the work, or the application of information presented in the manuscript, pose a threat to:

No	Yes
☐	☐ Public health
☐	☐ National security
☐	☐ Crops and/or livestock
☐	☐ Ecosystems
☐	☐ Any other significant area

Experiments of concern

Does the work involve any of these experiments of concern:

No	Yes
☐	☐ Demonstrate how to render a vaccine ineffective
☐	☐ Confer resistance to therapeutically useful antibiotics or antiviral agents
☐	☐ Enhance the virulence of a pathogen or render a nonpathogen virulent
☐	☐ Increase transmissibility of a pathogen
☐	☐ Alter the host range of a pathogen
☐	☐ Enable evasion of diagnostic/detection modalities
☐	☐ Enable the weaponization of a biological agent or toxin
☐	☐ Any other potentially harmful combination of experiments and agents

ChIP-seq

Data deposition

- ☐ Confirm that both raw and final processed data have been deposited in a public database such as [GEO](#).
- ☐ Confirm that you have deposited or provided access to graph files (e.g. BED files) for the called peaks.

Data access links

May remain private before publication.

For "Initial submission" or "Revised version" documents, provide reviewer access links. For your "Final submission" document, provide a link to the deposited data.

Files in database submission

Provide a list of all files available in the database submission.

Genome browser session

(e.g. [UCSC](#))

Provide a link to an anonymized genome browser session for "Initial submission" and "Revised version" documents only, to enable peer review. Write "no longer applicable" for "Final submission" documents.

Methodology

Replicates

Describe the experimental replicates, specifying number, type and replicate agreement.

Sequencing depth

Describe the sequencing depth for each experiment, providing the total number of reads, uniquely mapped reads, length of reads and whether they were paired- or single-end.

Antibodies

Describe the antibodies used for the ChIP-seq experiments; as applicable, provide supplier name, catalog number, clone name, and lot number.

Peak calling parameters

Specify the command line program and parameters used for read mapping and peak calling, including the ChIP, control and index files used.

Data quality

Describe the methods used to ensure data quality in full detail, including how many peaks are at FDR 5% and above 5-fold enrichment.

Software

Describe the software used to collect and analyze the ChIP-seq data. For custom code that has been deposited into a community repository, provide accession details.

Flow Cytometry

Plots

Confirm that:

- ☒ The axis labels state the marker and fluorochrome used (e.g. CD4-FITC).
- ☒ The axis scales are clearly visible. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers).
- ☒ All plots are contour plots with outliers or pseudocolor plots.
- ☒ A numerical value for number of cells or percentage (with statistics) is provided.

Methodology

Sample preparation

For isolation of splenic immune cells, the spleen was removed and mechanically disrupted, filtered through a 70-µm strainer, and red blood cells were lysed with 2 minutes incubation with ammonium chloride potassium buffer. Spleen cells were washed twice with cold PBS. For isolation of hepatic immune cells, murine liver was perfused via the portal vein for a minute and subsequently passed through a 100 µm mesh sieve, washed and the cell pellet was resuspended in 8 ml of GBSS containing collagenase IV (1:400, Worthington, #LS004189) and digested in a rotatory shaker (240 rpm) for 15 min at 37°C. The cell suspension was washed and lymphocytes were isolated through Percoll (40/80, GE Healthcare, #10607095) density centrifugation (7 acceleration, 1 deceleration, 1440?? g, for 25 min). For stimulation of lymphocytes, cells were stimulated with IL-15R?/IL-15 (10 ng/ml, ThermoFisher, #14-8152-80), IL-21 (10 ng/ml, Peprotech, # 210-21), IL-7 (10 ng/ml, Peprotech, # 217-17), IL-2 (10 ng/ml), TGF? (10 ng/ml, Peprotech, #100-21), IFN? (10 ng/ml, Peprotech, #315-05), IL-1 (10 ng/ml, Peprotech, #211-11B) or TNF (5 ng/ml, ThermoFisher, #PMC3013) for 18 hours followed by flow cytometry analysis. For isolation of lymphocytes from kidney, the renal capsula was removed and kidneys were digested by injecting 1 ml PBS containing type IV collagenase (1 mg/ml, Worthington, #LS004189) and DNase I (50 U/ml, Roche, #04716728001). After mechanical dissection, tissue was incubated for 60 min at 37°C, the cell suspension was filtered through a 70-µm mesh strainer, washed and subjected to Percoll (40%/80%) density centrifugation (1440 x g, for 25 min). For isolation of lymphocytes from murine skin, ears were cut and the dorsal side was removed from the ventral side. After cutting in small pieces, the tissue suspension was digested with 3 ml PBS solution containing type IV collagenase (2 mg/ml) and DNase I (50 U/ml) for 60 min at 37°C. Finally, the cell suspension was filtered through a 70-µm mesh strainer, washed and were ready for subsequent analysis. For isolation lymphocytes from the lung, both lobes were removed and digested by injecting 1 ml PBS solution containing type IV collagenase (1 mg/ml) and DNase I (50 U/ml). After tearing the lobes apart and removing remaining air by pipetting, tissue suspension was digested for 30 min at 37°C. Finally, tissue suspension was filtered through a 70-µm strainer, washed and subjected to Percoll (40%/80%)-density centrifugation (1440 x g, 25 min). To generate cell suspension from human liver tissue for flow cytometric analysis, tissue was mashed through a 100µm strainer and centrifuged (400 x g, 5 min). After cautiously decanting the supernatant, cell pellet was resuspended in medium (RPMI 1640 + 10% FCS + 1% P/S) and centrifuged again before proceeding with flow cytometric analysis.

PBMCs were generated from blood of healthy, voluntary donors which was approved by a vote from the ethics committee of the University Hospital München rechts der Isar. Blood was mixed with PBS (1:1) and loaded on Pancoll (PAN Biotech, #P04-601000) to isolate PBMCs through centrifugation (1400 g, 25 min).

Instrument

Sony SP6800 Spectral Analyser, Sony SA3800 Spectral Analyser, Sony SH800 Cell sorter

Software

FlowJo software, Tree Star, Ashland, USA v10

Cell population abundance

All samples were sorted using "high-purity" mode and every sample underwent a post-sort purity check. If sort purity was < 95.0% of the FSC/SSC gate of starting cell population

Gating strategy

FSC-A/SSC-A -> FSC-H/FSC-A -> FSC-A/Live-Dead -> cells of interest defined as "alive cells", followed by staining for T cells.

- ☒ Tick this box to confirm that a figure exemplifying the gating strategy is provided in the Supplementary Information.

Magnetic resonance imaging

Experimental design

Design type

Indicate task or resting state; event-related or block design.

Design specifications

Specify the number of blocks, trials or experimental units per session and/or subject, and specify the length of each trial or block (if trials are blocked) and interval between trials.

Behavioral performance measures

State number and/or type of variables recorded (e.g. correct button press, response time) and what statistics were used to establish that the subjects were performing the task as expected (e.g. mean, range, and/or standard deviation across subjects).

Acquisition

Imaging type(s)	<i>Specify: functional, structural, diffusion, perfusion.</i>
Field strength	<i>Specify in Tesla</i>
Sequence & imaging parameters	<i>Specify the pulse sequence type (gradient echo, spin echo, etc.), imaging type (EPI, spiral, etc.), field of view, matrix size, slice thickness, orientation and TE/TR/flip angle.</i>
Area of acquisition	<i>State whether a whole brain scan was used OR define the area of acquisition, describing how the region was determined.</i>
Diffusion MRI	☐ Used ☐ Not used

Preprocessing

Preprocessing software	<i>Provide detail on software version and revision number and on specific parameters (model/functions, brain extraction, segmentation, smoothing kernel size, etc.).</i>
Normalization	<i>If data were normalized/standardized, describe the approach(es): specify linear or non-linear and define image types used for transformation OR indicate that data were not normalized and explain rationale for lack of normalization.</i>
Normalization template	<i>Describe the template used for normalization/transformation, specifying subject space or group standardized space (e.g. original Talairach, MNI305, ICBM152) OR indicate that the data were not normalized.</i>
Noise and artifact removal	<i>Describe your procedure(s) for artifact and structured noise removal, specifying motion parameters, tissue signals and physiological signals (heart rate, respiration).</i>
Volume censoring	<i>Define your software and/or method and criteria for volume censoring, and state the extent of such censoring.</i>

Statistical modeling & inference

Model type and settings	<i>Specify type (mass univariate, multivariate, RSA, predictive, etc.) and describe essential details of the model at the first and second levels (e.g. fixed, random or mixed effects; drift or auto-correlation).</i>
Effect(s) tested	<i>Define precise effect in terms of the task or stimulus conditions instead of psychological concepts and indicate whether ANOVA or factorial designs were used.</i>
Specify type of analysis:	☐ Whole brain ☐ ROI-based ☐ Both
Statistic type for inference (See Eklund et al. 2016)	<i>Specify voxel-wise or cluster-wise and report all relevant parameters for cluster-wise methods.</i>
Correction	<i>Describe the type of correction and how it is obtained for multiple comparisons (e.g. FWE, FDR, permutation or Monte Carlo).</i>

Models & analysis

n/a	Involved in the study
☐	☐ Functional and/or effective connectivity
☐	☐ Graph analysis
☐	☐ Multivariate modeling or predictive analysis
Functional and/or effective connectivity	<i>Report the measures of dependence used and the model details (e.g. Pearson correlation, partial correlation, mutual information).</i>
Graph analysis	<i>Report the dependent variable and connectivity measure, specifying weighted graph or binarized graph, subject- or group-level, and the global and/or node summaries used (e.g. clustering coefficient, efficiency, etc.).</i>
Multivariate modeling and predictive analysis	<i>Specify independent variables, features extraction and dimension reduction, model, training and evaluation metrics.</i>