Human Intrahepatic Regulatory T Cells Are Functional, Require IL-2 From Effector Cells for Survival, and Are Susceptible to Fas Ligand-Mediated Apoptosis

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Regulatory T cells ($T_{reg}$) suppress T effector cell proliferation and maintain immune homeostasis. Autoimmune liver diseases persist despite high frequencies of $T_{reg}$ in the liver, suggesting that the local hepatic microenvironment might affect $T_{reg}$ stability, survival, and function. We hypothesized that interactions between $T_{reg}$ and endothelial cells during recruitment and then with epithelial cells within the liver affect $T_{reg}$ stability, survival, and function. To model this, we explored the function of $T_{reg}$ after migration through human hepatic sinusoidal-endothelium (postendothelial migrated $T_{reg}$ [PEM $T_{reg}$]) and the effect of subsequent interactions with cholangiocytes and local proinflammatory cytokines on survival and stability of $T_{reg}$. Our findings suggest that the intrahepatic microenvironment is highly enriched with proinflammatory cytokines but deficient in the $T_{reg}$ survival cytokine interleukin (IL)-2. Migration through endothelium into a model mimicking the inflamed liver microenvironment did not affect $T_{reg}$ stability; however, functional capacity was reduced. Furthermore, the addition of exogenous IL-2 enhanced PEM $T_{reg}$ phosphorylated STAT5 signaling compared with PEMCD8. CD4 and CD8 T cells are the main source of IL-2 in the inflamed liver. Liver-infiltrating $T_{reg}$ reside close to bile ducts and coculture with cholangiocytes or their supernatants induced preferential apoptosis of $T_{reg}$ compared with CD8 effector cells. $T_{reg}$ from diseased livers expressed high levels of CD95, and their apoptosis was inhibited by IL-2 or blockade of CD95.

Conclusion: Recruitment through endothelium does not impair $T_{reg}$ stability, but a proinflammatory microenvironment deficient in IL-2 leads to impaired function and increased susceptibility of $T_{reg}$ to epithelial cell-induced Fas-mediated apoptosis. These results provide a mechanism to explain $T_{reg}$ dysfunction in inflamed tissues and suggest that IL-2 supplementation, particularly if used in conjunction with $T_{reg}$ therapy, could restore immune homeostasis in inflammatory and autoimmune liver disease.

**Abbreviations:** ANOVA, analysis of variance; BEC, biliary epithelial cell; ELISA, enzyme-linked immunosorbent assay; FASL, Fas ligand; IL, interleukin; HSEC, hepatic sinusoidal endothelial cell; IFN-$\gamma$, interferon-$\gamma$; LIT$T_{reg}$, liver-infiltrating regulatory T cell; PEM$T_{reg}$, postendothelial migrated $T_{reg}$; SEM, standard error of the mean; Th 1, T helper 1; Th 17, T helper 17; TNF-$\alpha$, tumor necrosis factor-$\alpha$; $T_{reg}$, regulatory T cell.

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Intrahepatic T\textsubscript{reg} increase in parallel with effector T cells during chronic hepatitis\textsuperscript{(5)} demonstrates that the presence of T\textsubscript{reg} in the liver does not prevent ongoing hepatic inflammation. Whether this is due to dys-functional T\textsubscript{reg} or overwhelming effector responses is not known.

Understanding the fate of T\textsubscript{reg} in the inflamed liver is crucial, because T\textsubscript{reg} recruited to the liver from blood via hepatic sinusoids\textsuperscript{(6)} enter the hepatic microenvironment enriched with proinflammatory cytokines. The process of recruitment and/or exposure to this inflammatory microenvironment could affect T\textsubscript{reg} stability, survival, and function. Recruitment through endothelium can affect the differentiation and activation of various leukocyte subsets, and T\textsubscript{reg} can switch toward a T helper 1 (Th 1) or Th helper 17 (Th 17) lineage in an inflammatory environment.\textsuperscript{(7,8)} Furthermore, intrahepatic lymphocytes are removed by activation-induced cell death triggered by receptors such as CD95 (APO-1/Fas), leading to resolution of inflammation. Thus, differential susceptibility to apoptosis could affect the balance of T\textsubscript{reg} and T effector cells at inflammatory sites.\textsuperscript{(9,10)} T\textsubscript{reg} are highly sensitive to CD95-mediated apoptosis,\textsuperscript{(11)} and interleukin (IL)-2-dependent phosphorylation of STAT5 is crucial for their proliferation, differentiation, and survival.\textsuperscript{(12)} We reported previously that many intrahepatic T\textsubscript{reg} lack evidence of STAT5 phosphorylation, suggesting that the inflammatory liver microenvironment is hostile to T\textsubscript{reg} survival and function.\textsuperscript{(13)}

In the present study, we report that recruitment through endothelium into a model of the inflamed liver microenvironment reduces the suppressive capacity of T\textsubscript{reg} and a lack of local IL-2 enhances their susceptibility to Fas-mediated apoptosis induced by epithelial cells. These findings support the therapeutic potential of IL-2 therapy to restore local T\textsubscript{reg} function in inflammatory liver diseases.

Materials and Methods

Tissues and Blood

Venous blood was obtained from healthy volunteers or patients with hemochromatosis who were admitted to the Liver Unit at the Queen Elizabeth Hospital, Birmingham, UK\textsuperscript{(5)} (Local Research Ethics Committee reference no. 04/Q2708/41) and liver tissues from patients undergoing liver transplantation for inflammatory liver diseases, including primary biliary cirrhosis, primary sclerosing cholangitis, alcoholic liver disease, and autoimmune hepatitis. Normal liver was obtained from donor liver tissue surplus to clinical requirements (Local Research Ethics Committee reference no. 06/Q2708/11, 98/CA5192).

Isolation of Liver Infiltrating Lymphocytes, Biliary Epithelial Cells, and Primary Human Hepatic Sinusoidal Endothelial Cells

Liver-infiltrating lymphocytes,\textsuperscript{(5)} biliary epithelial cells (BECs),\textsuperscript{(14)} and hepatic sinusoidal endothelial cells (HSECs)\textsuperscript{(15)} were prepared and isolated from fresh liver tissue as described previously.\textsuperscript{(13)}

Postendothelial Transmigrated T\textsubscript{reg} and CD8\textsuperscript{+} T Cell Functional Assays

HSECs were plated and grown on six-well collagen I precoated Transwell inserts (3 μm; Greiner Bio-One) until confluent. Cells were then stimulated with 10 ng/mL interferon-γ (IFN-γ) and 10 ng/mL tumor necrosis factor α (TNF-α) (both Peprotech) for 24 hours to mimic the inflamed environment. After washing, the lower chamber
was filled with either Roswell Park Memorial Institute 1640 (RPMI-1640) medium (Gibco) supplemented with 0.05% bovine serum albumin or inflamed liver conditioned supernatant prepared by incubating 1 g of tissue in 4 mL serum-free RPMI 1640 for 24 hours. The supernatant was then centrifuged and filtered (0.22 μm-pore). Freshly isolated T_{reg} or CD8^+ T cells were then added to the upper chamber of the Transwell insert in 0.05% bovine serum albumin/RPMI-1640 and allowed to migrate across the endothelial cell monolayer into the lower chamber for 24 hours. The postendothelial transmigrated T_{reg} (PEM T_{reg}) or postendothelial transmigrated CD8^+ T (PEM CD8) cells were collected for experiments (T_{reg} suppression assay, biliary coculture assay, phosphorylated STAT5 assay, and T_{reg} plasticity assay).

PEM T_{reg} AND CD8 APOPTOSIS ASSAY

BECs were seeded into 24-well collagen I precoated plates, grown until confluent, and stimulated with 10 ng/mL IFN-γ and 10 ng/mL TNF-α (Peprotech) for 24 hours. PEM T_{reg} or PEM CD8 cells were cocultured with these BECs or with BEC supernatant for 24 hours, and apoptosis was analyzed by way of flow cytometric staining using Annexin and 7AAD (BD Pharmingen).

STATISTICAL ANALYSIS

Differences between treatments were evaluated by way of Student’s t test or one-way analysis of variance (ANOVA), followed by Bonferroni multiple comparison test using GraphPad Prism version 6.0 (GraphPad Software). P < 0.05 was considered statistically significant. Data are presented as the mean ± standard error of the mean (SEM).

See the Supporting Information for details regarding immunohistochemistry, confocal microscopy, enzyme-linked immunosorbent assay (ELISA), flow cytometry, isolation of peripheral blood T_{reg} and CD8 and assays of T_{reg} function, plasticity, and response to IL-2.

Results

FRESHLY ISOLATED HUMAN LIVER INFILTRATING T_{reg} ARE ACTIVATED, NONEXHAUSTED, MEMORY CELLS

We compared the frequency and surface phenotype of liver-infiltrating CD4^+CD25^{high}CD127^{low} T_{reg} (LIT_{reg}) in different inflammatory liver diseases with normal liver tissue (Fig. 1A,B) and with liver-infiltrating CD8 cells (Supporting Fig. S1C). T_{reg} isolated from normal and diseased livers differed in their expression of T_{reg}-associated functional surface receptors CD26, CD39, and CD69 (Fig. 1B). Very low expression of PD1 was observed on LIT_{reg} in both normal and diseased patients (9±5% versus 9.5±2%) (Fig. 1B) but high levels of CD44 were observed on LIT_{reg} (79±6% versus 87±5%) and liver-infiltrating CD8 cells (75±9% versus 90±4%) (Supporting Fig. S1C). LIT_{reg} exhibited a memory phenotype: CD45RO^{high}CD45RA^{low} and CCR7^{low} (85±7%) and inflamed liver tissue contained small populations of central memory CD45RA^{neg}CCR7^{pos} (4.7±3%) and tissue resident memory CD45RA^{pos}CCR7^{neg} (7±3%) LIT_{reg} (Fig. 1C,D).

PROINFLAMMATORY CYTOKINES ARE ELEVATED IN THE INFLAMED LIVER ENVIRONMENT, BUT THIS DOES NOT ALTER INTRAHEPATIC T_{reg} STABILITY

We explored the intrahepatic microenvironment by measuring proinflammatory cytokines in supernatants from inflamed human liver tissue. Diseased liver supernatants contained higher concentrations of IL-6 (8960 ± 4257 pg/mL), IL-8 (24,033 ± 16,589 pg/mL), IL-12 (61 ± 30 pg/mL), IFN-γ (32 ± 7 pg/mL), and IL-1β (363 ± 88 pg/mL) compared with normal liver supernatants (Fig. 2A).

We then examined the stability of intrahepatic T_{reg} in the hepatic microenvironment by modeling recruitment and intrahepatic conditions in vitro. Blood T_{reg} that had transmigrated across TNF-α and IFN-γ-stimulated HSECs had a phenotype similar to the LIT_{reg} phenotype, allowing us to use them to model the inflamed liver in vitro (Fig. 2B). PEM T_{reg} were migrated across stimulated HSECs into either control media or supernatants from inflamed liver tissue cultures (Supporting Fig. S5) and secretion of IL-10, IL-17, and IFN-γ by T_{reg} measured 24 hours postmigration (Fig. 2C) and expression of FOXP3, ROR, and Tbet analyzed up to 3 days (Fig. 2D, Supporting Fig. S2). A subset of LIT_{reg} expressed CD161 (20%) or IL-6 receptor (27%). We did not detect AhR expression (Fig. 2E).
PEM T\textsubscript{reg} REMAIN FUNCTIONALLY SUPPRESSIVE

After migration across endothelium into inflamed tissue supernatants, T\textsubscript{reg} maintained their ability to suppress the proliferation of T effector cells (Fig. 3A, Supporting Fig. S5A). Freshly isolated T\textsubscript{reg} were functionally suppressive (T\textsubscript{reg}/T effector cell ratio = 1:8) before any cell contact (Fig. 3Ai) and after contact with endothelium (T\textsubscript{reg}/T effector cell ratio = 1:4) but before transmigration (Fig. 3Aii). After transmigration across TNF-\textgamma and IFN-\gamma-stimulated HSECs into control media (Fig. 3Aiii), T\textsubscript{reg} had a reduced suppressive capacity at a 1:2 ratio. This was reduced further after migration into inflamed liver supernatant with suppression only seen at a 1:1 cell ratio (Fig. 3Aiv).

LIT\textsubscript{reg} expressed cell surface receptors associated with suppressive function (Fig. 3B,C) including CTLA-4 and CD147,\textsuperscript{(16)} whereas factors associated with cytolytic activity, granzyme-B, and perforin were expressed at higher levels on LICD8 T cells compared with LIT\textsubscript{reg} (53% versus 8% and 27% versus 8%). There were no significant differences in the expression of LAG-3 (21% versus 8%), CD137 (13% versus 4%), and TIM-3 (8% versus 4%) between LIT\textsubscript{reg} and LICD8 (Fig. 3B,C).

FIG. 1. Frequency and phenotype of intrahepatic T\textsubscript{reg} in inflamed human livers. Freshly isolated LIT\textsubscript{reg} from human explanted livers were phenotyped by flow cytometry. LIT\textsubscript{reg} were gated as CD4\textsuperscript{+}CD25\textsuperscript{high}CD127\textsuperscript{low}. (A) Frequency of LIT\textsubscript{reg} in normal livers and different diseased livers. (B) Surface marker expressions of LIT\textsubscript{reg} from normal livers, alcoholic liver disease livers, and autoimmune disease livers (see Supporting Fig. S1C for surface marker expressions of LICD8 cells from different diseased livers). LIT\textsubscript{reg} (defined as CD3\textsuperscript{+}CD4\textsuperscript{+}CD127\textsuperscript{low}CD25\textsuperscript{high}) were screened for expression of surface markers, CD26, CD39, CD44, CD69, and PD1. Data are presented as the mean \pm SEM (n = 6; one-way ANOVA followed by Bonferroni multiple comparison test). *P < 0.05. (C, D) Maturation status of LIT\textsubscript{reg}. Frequencies of CD45RA memory (CD45RA\textsuperscript{+}CCR7\textsuperscript{-}), CD45RO memory (CD45RO\textsuperscript{+}CCR7\textsuperscript{-}), and naive (CD45RA\textsuperscript{-}CCR7\textsuperscript{-}) LIT\textsubscript{reg} populations were determined by way of flow cytometry. Representative dot plots (C) and summary data (D) of LIT\textsubscript{reg} maturation status are shown. Data are presented as the mean \pm SEM (n = 6).
LIVER RESIDENT T_{reg} AND CD8 RESIDE AROUND Fas LIGAND (FASL)–POSITIVEBILE DUCTS

FOXP3\(^+\) LIT_{reg} were detected throughout the hepatic parenchyma, fibrous septa, and sinusoids as well as close to bile ducts in portal tracts (Fig. 4A–C) in inflamed liver tissue. CD95 ligand (FASL/CD178) was detected on CK19\(^+\) bile ducts in all inflammatory liver disease but not on normal liver tissue (Fig. 4D),\(^{14}\) with no difference in expression between different liver diseases.

CD95 EXPRESSION IS INCREASED ON LIVER INFILTRATING T_{reg} AND MEDIATES CELL APOPTOSIS IN RESPONSE TO BEC FASL

T_{reg} from diseased livers were more susceptible to apoptosis (Fig. 5A). LIT_{reg} expressed CD154, OX40, CD40, CD95, and CD27 and high levels of CD95 in all diseased livers (Fig. 5B). Liver-infiltrating CD8 cells were expressed at lower levels (Fig. 5C) Freshly isolated
Treg and CD8 cells were transmigrated across TNF-α and IFN-γ-stimulated hepatic sinusoidal endothelium into either control media, stimulated BEC supernatant, or contact with BECs (Fig. 6, Supporting Fig. S5B). Treg underwent apoptosis in this model, whereas very few CD8 T cells did. Treg apoptosis in response to contact with BECs or exposure to BEC supernatant was prevented by blocking CD95 ligand or by the addition of IL-2 (Fig. 6). Apoptotic immune cells were present around the bile duct, and we observed FOXP3 Treg and caspase-3 dual-positive apoptotic cells in peribiliary regions in tissue slides (Fig. 6C).

We detected high expression of CD27 on LITreg and CD70 the ligand for CD27 on liver-infiltrating dendritic
FIG. 4. Human diseased liver bile ducts expressing FASL and LITreg are present across liver lobules, with some residing in the peri-biliary region. Immunohistochemistry (A-D) and confocal fluorescence staining (Biii) of paraffin-embedded human liver sections for FOXP3, CD8, CK19 (biliary epithelial cell marker), and FASL are shown. (A) intrahepatic FOXP3+ Treg are present across the parenchyma (i), septa (ii), and portal tract (iii). (B) LICD8+ T cells reside close to bile ducts in the portal tracts in diseased human liver sections (Bi) alcoholic liver disease (ALD), (Bii) primary biliary cirrhosis (PBC), (Biii) Localization of CD8 cells on bile ducts in AIH liver tissue (red = CD8 [phycoerythrin]; green = bile ducts/CK19 [fluorescein isothiocyanate]; blue = nuclear stain [4',6-diamidino-2-phenylindole]). (C) Localization of FOXP3+ cells around the bile duct in diseased livers (Ci) autoimmune hepatitis (AIH), (Cii) hepatitis C virus (HCV), (Ciii) non A non B seronegative hepatitis. (D) Expression of FASL (right panel) by human CK19-expressing bile ducts (left panel) in normal and diseased livers.
cells. However, recombinant CD70 had no effect on Treg or CD8 T cell proliferation (Supporting Fig. S4).

**EFFECTOR T CELLS ARE THE MAIN SOURCE OF INTRAHEPATIC IL-2 AND T\textsubscript{reg} RESPOND BY PHOSPHORYLATION OF STAT5**

IL-2 was nearly undetectable in supernatants prepared from normal and chronic liver disease tissues (Fig. 7A). Secretion of IL-2 by human hepatocytes, HSECs, stromal cells, and biliary epithelial cells was minimal (Fig. 7B). IL-2 expression by liver-infiltrating activated CD4/CD8 cells was observed by way of flow cytometry (Fig. 7C) and secreted IL-2 was detected in the supernatant of liver-infiltrating CD4 and CD8 lymphocytes after stimulation with anti-CD3/CD28 beads, suggesting that effector lymphocytes are the main source of IL-2 in inflamed livers (Fig. 7D).

A high level of STAT5 phosphorylation was induced in 80%-90% of T_{reg} both non-transmigrated and PEM T_{reg} (Fig. 8, Supporting Fig. S4), and was not affected by transmigration into the inflamed liver microenvironment (Fig. 8). Significantly fewer CD8 cells (27%-36%) responded compared with T_{reg} (Supporting Fig. S4), and the level of phosphorylated STAT5 generated was also significantly greater in T_{reg} than in CD8 T cells (Fig. 8). Thus, IL-2-CD25 signaling and STAT5 phosphorylation remained intact in T_{reg} after transmigration into the inflamed hepatic microenvironment.

**Discussion**

Functionally active T_{reg} are required to suppress effector cells and maintain immune homeostasis, and the balance of T_{reg} and T effector cells will contribute to the outcome of liver inflammation. Therapeutic modulation of tissue resident T_{reg} has potential for treating autoimmune liver diseases. Once T_{reg} have migrated across the hepatic sinusoid, they are exposed to cytokines including IL-1\textbeta, IL-6, IL-8, IL-12, and IFN-\gamma in the inflamed hepatic microenvironment (Fig. 2). This proinflammatory milieu has the potential to influence the stability and function of intrahepatic T_{reg}. We observed that PEM T_{reg} that had migrated through activated endothelium into liver tissue supernatants showed no changes in FOXP3, RORc, or Tbet expression at 3 days, suggesting that T_{reg} differentiation is...
not affected by the liver milieu. In particular, the lack of ligand-activated transcription factor aryl hydrocarbon receptor and low levels of CD161, which are associated with Th 17 polarity, suggest few cells differentiate into Th 17 cells within the inflamed environment.\(^{21-23}\)

The frequency of LIT\(_{\text{reg}}\) is increased across all liver diseases,\(^{24}\) and these cells are not exhausted, with very low expression of PD-1 and high levels of CD69 and maintained functional properties\(^{25,26}\) (Fig. 3Aiv). LIT\(_{\text{reg}}\) exposed to a liver microenvironment maintain high levels of CD39, an ectonucleotidase used by T\(_{\text{reg}}\) to generate

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**FIG. 6.** Apoptosis of PEM T\(_{\text{reg}}\) but not CD8 is FASL dependent and can be rescued by blocking FASL or by supplementing with IL-2. T\(_{\text{reg}}\) and CD8+ T cells were transmigrated across TNF-\(\alpha\)- and IFN-\(\gamma\)-stimulated human sinusoidal endothelium to obtain postendothelial migrated cells, which were then cocultured with BECs or BEC supernatant in the presence or absence of recombinant IL-2 or anti-FASL antibody. Apoptosis was analyzed using flow cytometry for Annexin V. (A) Representative dot plots of apoptosis for T\(_{\text{reg}}\) and CD8 before and after endothelial migration. (B, C) Apoptosis of PEM T\(_{\text{reg}}\) (B) and PEM CD8 cells (C) with either BECs or BEC supernatant in the presence or absence of IL-2 or anti-FASL (one-way ANOVA followed by Bonferroni multiple comparison tests). **P ≤ 0.05. (D) Terminal deoxynucleotidyl transferase-mediated deoxyuridine triphosphate nick-end labeling revealed that apoptotic immune cells were present around bile ducts. (E) Dual staining of FOXP3 (brown) and caspase-3 (blue) in different diseased livers.
Immunosuppressive adenosine from extracellular nucleotides.\(^{27}\) Low expression of CD39 has been reported in autoimmune liver disease, and our data suggest that the hepatic microenvironment cannot explain this finding.\(^{28}\)

Similarly, LIT\(_{\text{reg}}\) expressed high levels of CD44, which has been associated with FOXP3 expression and suppressive function.\(^{29}\) After transendothelial migration into inflamed liver supernatant, T\(_{\text{reg}}\) maintained IL-10
secretion and CD147 expression, a marker of activated and suppressive T_reg. In addition, we also observed an enrichment of CTLA-4 expression in LIT_reg. CTLA-4 interacts with and removes CD80/CD86 from dendritic cells by way of transendocytosis, resulting in impaired costimulation via CD28 and is thus critical for T_reg function. Overall, LIT_reg are equipped with key surface markers and intracellular cytokines required to execute their suppressive capacity. These findings are encouraging for proposed studies of adoptive T_reg therapy in autoimmune and inflammatory liver diseases because they suggest that therapeutic cells will remain functional and will not differentiate into effector cells in an inflammatory environment.

We detected LIT_reg and LICD8 cells throughout the parenchyma and fibrous septa as well as around bile ducts in patients with liver disease. Biliary epithelial cells can secrete chemokines that localize...
lymphocytes to portal tracts. We recently reported that VCAM-1 on bile ducts supports effector T cell survival by binding to α4β1, but apoptosis pathways affecting different lymphocyte subsets in the inflamed liver are poorly understood. BECs express FASL, which can induce T cell death by activating Fas. Thus, expression or secretion of FASL by BECs could contribute to intrahepatic T cell apoptosis.

Circulating T$_\text{reg}$ are highly susceptible to CD95-FASL-dependent apoptosis but not to TCR-mediated cell death in contrast to effector T cells. T$_\text{reg}$ in the inflamed liver undergo CD95-dependent cell death as a consequence of FASL expression by tumor cells. We have reported previously that BECs express FASL in inflammatory liver diseases, but we have not explored the role of Fas in the differential fate of bile duct-associated T$_\text{reg}$ and CD8 cells. In the present study, we demonstrate that PEM T$_\text{reg}$ are more susceptible to CD95-FASL-mediated apoptosis either in contact with BECs or in response to BEC supernatant compared with PEM CD8 cells. T$_\text{reg}$ could be rescued by blocking FASL on BECs or in the inflamed biliary supernatant. If apoptosis of T$_\text{reg}$ affects only those in contact with bile ducts, and not those elsewhere in the liver, this may not have a substantial effect on the overall frequency of T$_\text{reg}$ in the diseased liver. In addition, the total frequency of T$_\text{reg}$ in the inflamed liver will depend on other factors, including the balance between recruitment, retention, proliferation, and exit of cells from the liver.

IL-2 is a crucial cytokine for T$_\text{reg}$ survival that activates STAT5 phosphorylation in response to activation of the IL-2 receptor. IL-2 can selectively expand T$_\text{reg}$ populations in vitro and in vivo. We detected minimal IL-2 in both the normal and inflamed liver microenvironment, which might account in part for reduced T$_\text{reg}$ functional capacity in the inflammatory tissue. The intrahepatic source of IL-2 is activated in CD4 and CD8 cells, which may be rapidly consumed by both effector and T$_\text{reg}$ for their survival and proliferation. Exogenous IL-2 restored levels of STAT5 phosphorylation and protected T$_\text{reg}$ from Fas-mediated apoptosis but had no effect on CD8 T cells. Thus, the lack of IL-2 in the inflamed liver microenvironment may result in upregulated Fas-mediated T$_\text{reg}$ apoptosis as well as contributing to defective intrahepatic LIT$_\text{reg}$ function. Because the requirement for IL-2 is much higher for CD8 cells, they may depend on different survival mechanisms. This is in agreement with a previous study on the effect of IL-2 on peripheral blood lymphocyte subsets. This enhanced phosphorylation of STAT5 in T$_\text{reg}$ compared with CD8 cells in response to IL-2 occurred in the presence and absence of cytokine-enriched inflamed hepatic supernatant, suggesting that the differential effect of IL-2 on T$_\text{reg}$ compared with effector cells is maintained in an inflammatory environment. This finding provides support for the therapeutic manipulation of LIT$_\text{reg}$ by treating these patients with IL-2 along with T$_\text{reg}$ therapy in the future to maintain functional and surviving LIT$_\text{reg}$.

In conclusion, we show for the first time that LIT$_\text{reg}$ are phenotypically stable within the inflamed hepatic environment and exhibit minimal plasticity. They remain functionally suppressive but are susceptible to apoptosis via the Fas pathway mediated by FASL on bile ducts, and this can be inhibited by exogenous IL-2. Therefore, IL-2 therapy may successfully restore intrahepatic T$_\text{reg}$ numbers and function in liver disease.

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Supporting Information

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