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Lkb1 maintains T_{reg} cell lineage identity

Di Wu^{1,*}, Yuechen Luo^{1,*}, Wei Guo^{1,*}, Qing Niu¹, Ting Xue², Fei Yang¹, Xiaolei Sun¹, Song Chen¹, Yuanyuan Liu¹, Jingru Liu³, Zhina Sun¹, Chunxiao Zhao¹, Huifang Huang³, Fang Liao², Zhongchao Han¹, Dongming Zhou⁴, Yongguang Yang⁵, Guogang Xu⁶, Tao Cheng¹ & Xiaoming Feng¹

Regulatory T (T_{reg}) cells are a distinct T-cell lineage characterized by sustained *Foxp3* expression and potent suppressor function, but the upstream dominant factors that preserve T_{reg} lineage-specific features are mostly unknown. Here, we show that Lkb1 maintains T_{reg} cell lineage identity by stabilizing *Foxp3* expression and enforcing suppressor function. Upon T-cell receptor (TCR) stimulation Lkb1 protein expression is upregulated in T_{reg} cells but not in conventional T cells. Mice with T_{reg} cell-specific deletion of Lkb1 develop a fatal early-onset autoimmune disease, with no *Foxp3* expression in most T_{reg} cells. Lkb1 stabilizes *Foxp3* expression by preventing STAT4-mediated methylation of the conserved noncoding sequence 2 (CNS2) in the *Foxp3* locus. Independent of maintaining *Foxp3* expression, Lkb1 programs the expression of a wide spectrum of immunosuppressive genes, through mechanisms involving the augmentation of TGF- β signalling. These findings identify a critical function of Lkb1 in maintaining T_{reg} cell lineage identity.

¹State Key Laboratory of Experimental Hematology, Institute of Hematology and Hospital of Blood Diseases, Chinese Academy of Medical Sciences & Peking Union Medical College, Tianjin 300020, China. ²Department of Medical Microbiology, Tongji Medical College, Huazhong University of Science and Technology, Wuhan 430030, China. ³Central Laboratory, The Union Hospital of Fujian Medical University, 29 Xinquan Road, Fuzhou 350001, China. ⁴Institute Pasteur of Shanghai, Shanghai Institutes for Biological Sciences, Chinese Academy of Sciences, Shanghai 200031, China. ⁵First Hospital of Jilin University, Changchun 130012, China. ⁶Nanlou Respiratory Department, Chinese PLA General Hospital, Beijing 100853, China. *These authors contributed equally to this work. Correspondence and requests for materials should be addressed to X.F. (email: fengxiaoming@ihcams.ac.cn) or to T.C. (email: chengtao@ihcams.ac.cn).

Regulatory T (T_{reg}) cells preserve immune homeostasis by suppressing autoreactive immune responses^{1,2}. The T_{reg} cell lineage can be defined by two basic characteristics, stable expression of the transcription factor Foxp3 (forkhead box P3) and potent suppressive capacity^{3,4}. T_{reg} cells are stable and usually retain lineage characteristics *in vivo*^{5,6}. However, under some conditions, T_{reg} cells can lose or alter their lineage identity, resulting in immune disturbance and development of diseases^{3,4,6,7}. Thus, delineating the molecular mechanisms that maintain the T_{reg} lineage identity is important for understanding and treating T_{reg} cell-related immune diseases.

Foxp3 is the most specific marker for distinguishing T_{reg} cells from other T cells and an important regulator required for programming T_{reg} suppressive function^{1,2}. Lineage tracing in mice has shown that thymus-derived mature T_{reg} cells have relatively stable Foxp3 expression in both homeostasis and inflammatory conditions⁵. Other reports indicate that epigenetic demethylation of the conserved noncoding sequence 2 (CNS2, also known as T_{reg} cell-specific demethylation region) in the *Foxp3* locus can ensure stable Foxp3 expression in T_{reg} cells^{8–10}. However, the upstream signalling checkpoints that activate chromatin, and the demethylation status, of *Foxp3* locus are not clear.

T_{reg} cells suppress immune responses through diverse mechanisms, such as the modulation of antigen presentation function (via CTLA4 (cytotoxic T-lymphocyte associated protein 4)), the production of inhibitory cytokines (for example, interleukin (IL)-10 and IL-35) and metabolites (for example, reactive oxygen species and adenosine), the deprivation of the T-cell growth factor IL-2 (via CD25), and the direct killing of target cells (via granzyme B and perforin)^{2,11,12}. Although Foxp3 has an important function in programming T_{reg} cells by controlling the expression of a large number of immunosuppressive genes, Foxp3 alone is not sufficient to confer this function^{1,2}. In addition, T-cell receptor (TCR) signalling is also important to promote T_{reg} cell function^{13–15}. Nevertheless, little is known about other upstream ‘master regulators’ that broadly control the expression of these T_{reg} cell-associated immunosuppressive genes.

Liver kinase b1 (Lkb1) is a tumour suppressor, and is mutated in Peutz–Jeghers cancer syndrome, cervical carcinoma and many sporadic non-small-lung carcinomas^{16–18}. Under energy-stressed conditions, Lkb1 is an important upstream kinase that phosphorylates AMP-activated protein kinase (AMPK) and AMPK-related kinases that coordinate cell growth with metabolism^{16–18}. Lkb1 has been shown to restrain the activation and proinflammatory function of conventional T cells^{18,19}.

In this study, we find that Lkb1 protein is specifically increased in T_{reg} cells upon TCR stimulation. To understand the function and mechanism of Lkb1 in T_{reg} cells, we generate a mouse line with Lkb1 specifically deleted in T_{reg} cells. These mice develop a fatal early-onset autoimmune disease with defective maintenance of stable *Foxp3* expression and suppressive capacity in T_{reg} cells. Mechanistically, Lkb1 restrains STAT4 (signal transducer and activator of transcription 4) activation partially through suppressing nuclear factor- κ B (NF- κ B) signalling, and thus prevents STAT4-mediated methylation of CNS2 in the *Foxp3* locus, resulting in stable Foxp3 expression. Meanwhile, Lkb1 promotes the expression of a large number of immunosuppressive genes partially through augmenting transforming growth factor- β (TGF- β) signalling. Our study identifies Lkb1 as a critical determinant of T_{reg} cell lineage identity.

Results

Lkb1 protein is increased in T_{reg} cells upon TCR stimulation. TCR stimulation is essential for T_{reg} cells to exert their optimal

function^{13–15}. Lkb1 protein expression in T_{reg} cells were slightly lower compared with conventional T cells without stimulation (Fig. 1a and Supplementary Figs 1a and 8a,b). However, upon TCR stimulation, Lkb1 protein expression was markedly upregulated in T_{reg} but not conventional T cells (Fig. 1a and Supplementary Fig. 1a), implying that Lkb1 might be particularly important for T_{reg} cells to execute their immunoregulatory effect.

Deletion of Lkb1 in T_{reg} cells leads to a fatal autoimmunity.

To investigate the role of Lkb1 in T_{reg} cells, we generated a mouse line with Lkb1 conditionally depleted in T_{reg} cells by crossing *Foxp3*^{YFP-Cre} (*Foxp3*^{Cre}, which ensure T_{reg} cell-specific deletion of a target gene) mice²⁰ with *Lkb1*^{fl/fl} mice²¹. Lkb1 protein was depleted in CD4⁺YFP⁺ T_{reg} cells from *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice (Supplementary Figs 1b and 8c,d). Strikingly, loss of Lkb1 in T_{reg} cells caused early moribund at ~35 days of age (Fig. 1b). *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice exhibited smaller size, decreased mobility, hunched posture, collapsed ears and tail skin lesions (Fig. 1c), and barely increased in body weight with age (Fig. 1d). *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice displayed splenomegaly and lymphadenopathy (Fig. 1e), and they had increased cell numbers in secondary lymphoid organs (Fig. 1f). Histopathological analysis revealed massive infiltration of lymphocytes into multiple organs such as skin, lung, liver and stomach (Fig. 1g), suggesting T-cell autoimmunity. In support of this, *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice had substantial expansion of CD4⁺ conventional T cells (Supplementary Fig. 1c), associated with drastically increased percentages of cells that displayed a CD44^{high}CD62L^{low} effector/memory phenotype (Fig. 2a and Supplementary Fig. 1d). These mice also had higher percentages of CD4⁺ and CD8⁺ conventional T cells that expressed the proliferation marker Ki67 and acute activation markers CD25 and CD69 (Fig. 2b and Supplementary Fig. 1e), and produced the inflammatory cytokines interferon- γ , IL-4 and IL-17A (Fig. 2c and Supplementary Fig. 1f). These phenotypes were similar in severity to those observed in mice deficient of Foxp3 (ref. 22) or depleted of T_{reg} cells²³, indicating a severe defect of immune suppression mediated by T_{reg} cells.

Lkb1 maintains *Foxp3* expression and CNS2 demethylation.

T_{reg} cells normally stay at a relatively constant abundance within the CD4⁺ T-cell population, and expand concomitantly with effector/memory T cells to preserve immune homeostasis during inflammation²⁴. Although not altered at early ages, the percentages of peripheral T_{reg} cells among CD4⁺ T-cell populations continuously dropped to ~3% by 4 weeks of age when the mice became moribund (Fig. 3a). Despite greatly increased numbers of effector/memory T cells in *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice (Supplementary Fig. 1d), the absolute numbers of T_{reg} cells were only comparable with those in wild-type mice (Supplementary Fig. 1g), suggesting that Lkb1-deficient T_{reg} cells fail to accumulate concomitantly with effector/memory T cells in *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice at later ages that might contribute to the autoimmunity in *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice.

The survival and proliferation of Lkb1-deficient T_{reg} cells were barely altered compared with wild-type control cells (Supplementary Fig. 2a,b). Thus, other reasons might account for the defective accumulation of Lkb1-deficient T_{reg} cells. T_{reg} cells are considered a stable lineage marked by sustained Foxp3 expression⁵, but under certain conditions they might lose Foxp3 expression and develop into cells resembling effector T cells (designated ex- T_{reg} cells)²⁵. To investigate whether Lkb1-deficient T_{reg} cells were prone to lose Foxp3 expression, we introduced the *Rosa26*^{YFP} allele (a loxp-site-flanked STOP cassette followed by the YFP-encoding sequence was inserted into the *Rosa26* locus, and the expression of YFP from the *Rosa26*

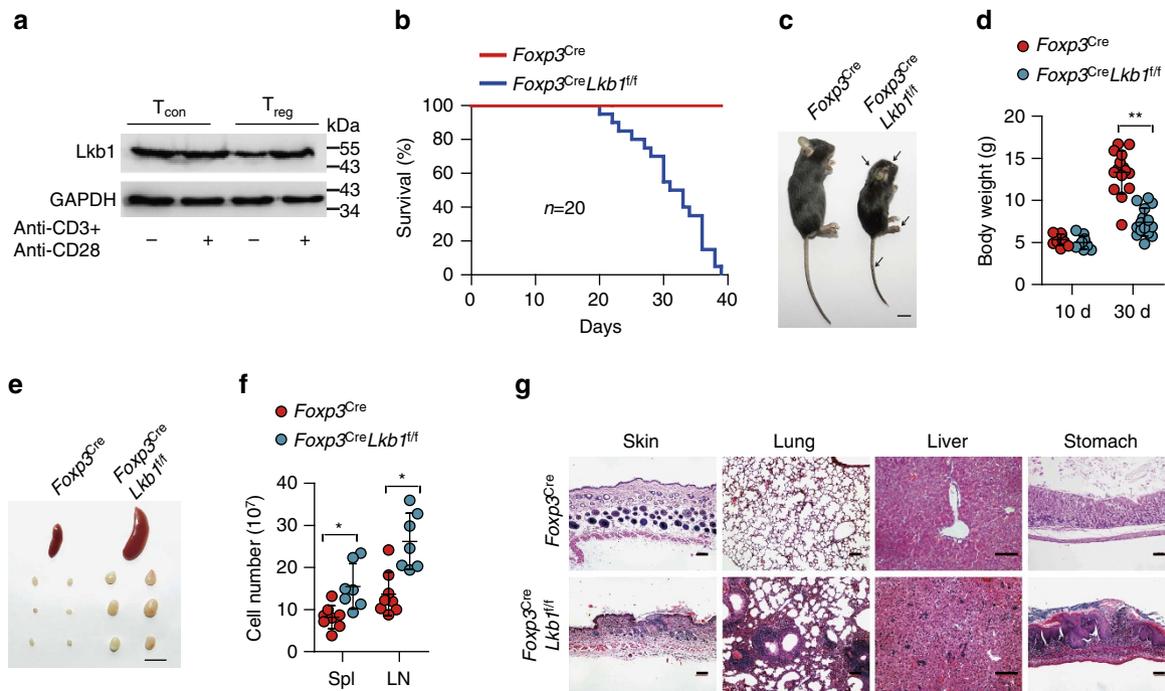


Figure 1 | *T_{reg}* cell-specific deletion of *Lkb1* leads to a fatal autoimmune disease. (a) *Lkb1* proteins in $CD4^{+} YFP^{-}$ conventional T cells and $CD4^{+} YFP^{+}$ *T_{reg}* cells untreated or stimulated in plates coated with anti-CD3 and anti-CD28 in the presence of IL-2 for 24 h determined by western blot. (b) Survival of *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice ($n=20$). (c) A representative appearance of 30-day-old *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice (scale bar, 1 cm). (d) Body weight of *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* male mice of different ages ($n=10-15$). (e) A representative picture of spleen and lymph nodes from 30-day-old *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice. (scale bar, 1 cm). (f) Total cell numbers in spleen (Spl) and lymph nodes (LNs) of *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice ($n=7-8$). (g) Representative haematoxylin and eosin-stained skin, lung, liver and stomach sections from *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice (scale bar, 100 μ m). All mice analysed were 28–30 days old, unless otherwise specified. Log-rank survival curve was used for survival analysis in **b**, and two-way analysis of variance (ANOVA) was used for statistical analyses in **d,f** ($*P<0.05$, $**P<0.01$, $****P<0.0001$); error bars represent s.d.; all data are representative of at least three independent experiments.

locus was dependent on the expression of Cre recombinase) into wild-type and conditional knockout mice²⁶. $CD4^{+} Rosa26-YFP^{high}$ cells (100% *Lkb1*-deficient *T_{reg}* cells) sorted from *Foxp3^{Cre}Lkb1^{fl/fl}Rosa26^{YFP}* mice contain much higher percentages of *Foxp3⁻* cells (which represented ex-*T_{reg}* cells that ever experienced *Foxp3* expression but ceased to express it later) than that from *Foxp3^{Cre}Rosa26^{YFP}* mice (Fig. 3b). Therefore, a large proportion of *Lkb1*-deficient *T_{reg}* cells lost *Foxp3* expression and became ex-*T_{reg}* cells that might result in the decrease of *T_{reg}* cell abundance *in vivo*.

Foxp3 is an X-chromosome-encoded transcription factor. Due to random X inactivation, female mice heterozygous for *Foxp3^{Cre}* (*Foxp3^{Cre/+}*) should contain ~1:1 ratio of *Foxp3^{Cre}*-positive *T_{reg}* cells to *Foxp3^{Cre}*-negative *T_{reg}* cells. Hence, the *Foxp3^{Cre}*-heterozygous female *Foxp3^{Cre/+}Lkb1^{fl/fl}* mice were devoid of autoimmune diseases due to the presence of *T_{reg}* cells lacking *Foxp3^{Cre}* expression but retaining a wild-type *Lkb1* allele⁸. Proinflammatory cytokines may impair *T_{reg}* cell stability^{3,6}. To determine whether the instability of *T_{reg}* cells was due to either the inflammatory environment in *Foxp3^{Cre}Lkb1^{fl/fl}Rosa26^{YFP}* mice or the intrinsic defect of the cells, we utilized *Foxp3^{Cre}*-heterozygous female *Foxp3^{Cre/+}Lkb1^{fl/fl}Rosa26^{YFP}* mice that were devoid of autoimmune diseases. Despite the presence of wild-type *T_{reg}* cells devoid of the expression of *Foxp3^{Cre}* in *Foxp3^{Cre/+}Lkb1^{fl/fl}Rosa26^{YFP}* mice, the $CD4^{+} Rosa26-YFP^{high}$ cells from these mice were cells that had ever experienced the expression of *Foxp3^{Cre}* that resulted in the deletion of *Lkb1* and expression of *Rosa26-YFP*. Similarly, we observed substantially higher proportions of *Foxp3⁻* cells among sorted

$CD4^{+} Rosa26-YFP^{high}$ cells from *Foxp3^{Cre/+}Lkb1^{fl/fl}Rosa26^{YFP}* mice than that from *Foxp3^{Cre/+}Rosa26^{YFP}* mice (Supplementary Fig. 2c), ruling out the possibility that overproduction of proinflammatory cytokines in *Foxp3^{Cre}Lkb1^{fl/fl}* mice were the causative reasons for *T_{reg}* cell instability.

Foxp3⁻Rosa26-YFP⁺ ex-*T_{reg}* cells might also arise from recently activated T cells that transiently expressed *Foxp3* but did not develop into a full *T_{reg}* program (which were also considered as poorly committed *T_{reg}* cells)²⁷. To determine whether the full committed *T_{reg}* cells lose stability, we doubly sorted $CD4^{+} CD25^{+} YFP^{+}$ cells from $CD45.1^{+} CD45.2^{+}$ wild-type *Foxp3^{Cre}* and $CD45.1^{-} CD45.2^{+} Foxp3^{Cre}Lkb1^{fl/fl}$ mice, mixed them together with congenitally marked $CD45.1^{+} CD45.2^{-}$ naive T (T_n) cells and transferred them into *Rag1^{-/-}* mice. Strikingly, we observed that nearly all the *Lkb1*-deficient *T_{reg}* cells lost *Foxp3* expression 3 weeks after transfer (Fig. 3c), despite the fact that most wild-type *T_{reg}* cells retained *Foxp3* expression. This was not due to the contamination of non-*T_{reg}* cells because the starting population contained >99.5% *Foxp3⁺* *T_{reg}* cells after double sorting (Supplementary Fig. 2d). These results confirm that *Foxp3* expression is lost in a large proportion of fully committed mature *Lkb1*-deficient *T_{reg}* cells, rather than in a subset of poorly committed *Foxp3⁺* cells.

Next, we investigated the molecular mechanisms by which *Lkb1* promoted *T_{reg}* cell stability. Because *Foxp3* protein expression level was barely altered in *Lkb1*-deficient *T_{reg}* cells (Supplementary Fig. 2e), *Lkb1* seemed to not control *T_{reg}* stability through affecting *Foxp3* protein production or degradation. The epigenetic status across the *Foxp3* locus has been shown to be

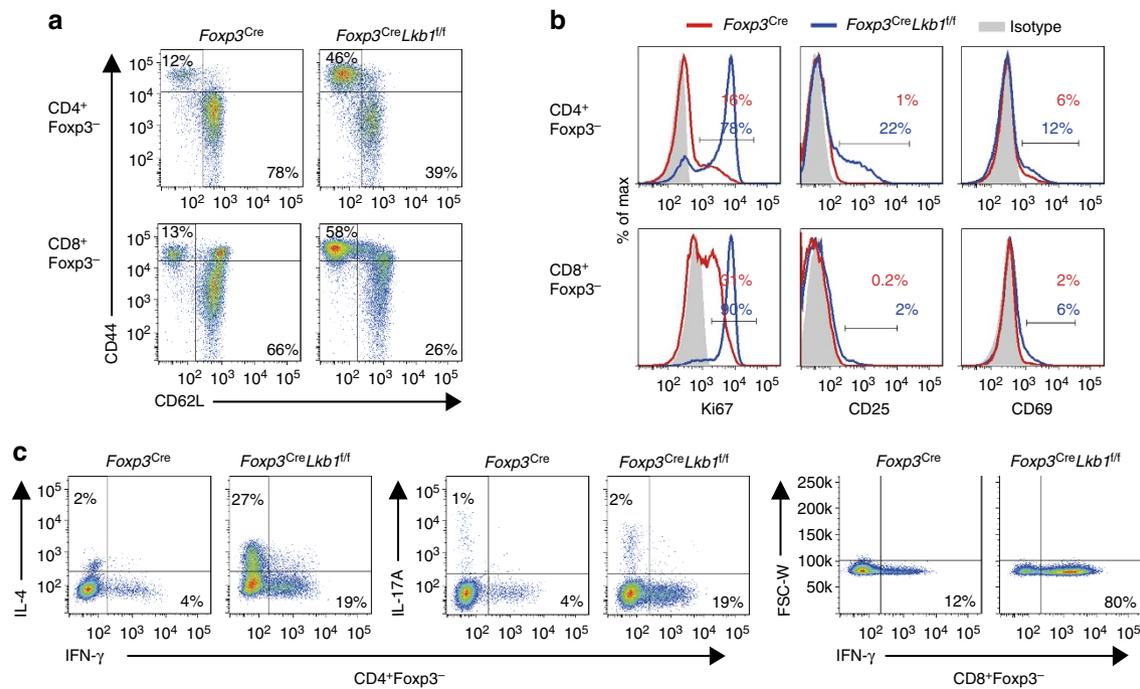


Figure 2 | Spontaneous activation of conventional T cells in *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice. (a) Expression of CD44 and CD62L on splenic CD4⁺ Foxp3⁻ and CD8⁺ Foxp3⁻ T cells from *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice. **(b)** Expression of Ki67, CD25 and CD69 in splenic CD4⁺ Foxp3⁻ and CD8⁺ Foxp3⁻ T cells from *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice. **(c)** Intracellular staining of cytokines in splenic CD4⁺ Foxp3⁻ and CD8⁺ Foxp3⁻ T cells from *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice stimulated with phorbol myristate acetate (PMA) and ionomycin for 4 h. All mice analysed were 28–30 days old. All data are representative of at least three independent experiments.

critical for sustained Foxp3 expression⁹. Two recent reports demonstrated that deletion of CNS2 region in the *Foxp3* locus led to the loss of T_{reg} cell stability^{8,10}, suggesting that CNS2 is pivotal for maintaining stable Foxp3 expression in T_{reg} cells. CNS2 is completely demethylated in T_{reg} cells, but not in conventional T cells^{9,28–30}. T_{reg} cell lineage development is partially dependent on the demethylation of CNS2 (refs 31,32), and CNS2 has enhancer activity that is markedly reduced after methylation³¹. Therefore, we examined the methylation of CpG motifs across the *Foxp3* locus by bisulfite sequencing in CD4⁺ YFP⁺ T_{reg} cells from *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice and *Foxp3*^{Cre} mice, and found a significant increase (nearly 25%) in methylation of CpG at CNS2 in *Lkb1*-deficient T_{reg} cells *ex vivo* (Fig. 3d). T_{reg} cells that already lost Foxp3 expression were excluded in this analysis. Although *Foxp3* promoter and other potentially regulatory regions are associated with T_{reg} cell lineage identity^{29,33,34}, the methylation of CpG at sites including –1.5 kb region, promoter, exon 1, exon 8 and exon 11 across the *Foxp3* locus in *Lkb1*-deficient T_{reg} cells was comparable to the wild-type T_{reg} cells (Fig. 3d). Therefore, we hypothesize that methylation of CpG at CNS2 may contribute to the instability of *Lkb1*-deficient T_{reg} cells. It seems that the methylation of CNS2 is not a secondary effect of Foxp3 reduction, because Foxp3 protein level was barely altered in *Lkb1*-deficient T_{reg} cells that already had substantial CNS2 methylation. Given the critical role of CNS2 in conferring T_{reg} cell stability established by previous studies, our results together suggested that CNS2 methylation might lead to the instability of *Lkb1*-deficient T_{reg} cells.

Lkb1 functions in T_{reg} cells independent of AMPK. AMPK is the well-known *Lkb1* downstream target critical for coordinating metabolism¹³. Interestingly, we found no obvious decrease of AMPK activity in *Lkb1*-deficient T_{reg} cells (Supplementary Figs 3a and 8e–h). Therefore, other kinases such as

calmodulin-dependent protein kinase-β (CamKKβ) and transforming growth factor-β-activated kinase-1 (TAK1)¹⁵ may promote AMPK activation in T_{reg} cells. These data argue that *Lkb1* is not important for AMPK activation in T_{reg} cells, and *Lkb1* substrates other than AMPK are responsible for the observed phenotype of *Lkb1*-deficient T_{reg} cells. Consistent with this, T_{reg} cells that lacked the expression of AMPKα1/2 (from *Foxp3*^{Cre}*AMPKα1*^{fl/fl}*AMPKα2*^{fl/fl} mice, Supplementary Fig. 3b) did not exhibit any adverse phenotype and *in vivo* functional impairment (Supplementary Fig. 3c,d).

Lkb1 prevents STAT4 activation and STAT4 binding to CNS2. STAT3 and STAT6 are respectively the downstream mediators of the inflammatory cytokines IL-6 and IL-4 that bind to *Foxp3* gene to mediate CpG methylation and T_{reg} cell instability⁸, whereas the T_{reg} cell growth factor IL-2 signals via STAT5 to counteract the destabilizing effect of these inflammatory cytokines⁸. Therefore, we examined whether the STAT family protein activations were altered in *Lkb1*-deficient T_{reg} cells. Although STAT3 and STAT5 phosphorylations were barely affected, we detected a drastic increase of STAT4 phosphorylation and a slight increase of STAT6 phosphorylation in *Lkb1*-deficient T_{reg} cells with or without the stimulation of relevant cytokines (Fig. 4a and Supplementary Figs 4a,b and 8i–k). The *in vivo* maintenance of T_{reg} cells depends on their contact with dendritic cells (DCs)^{35,36}, and hence we developed an *in vitro* DC-T_{reg} cell co-culture system that could mimic the physiological situation to test which signalling would cause the instability of T_{reg} cells. *Lkb1*-deficient T_{reg} cells slightly lost stability when co-cultured with DCs without exogenous cytokines (Fig. 4b). Treatment with IL-2 alone, or IL-2 in combination with IL-4 or IL-6, did not lead to a significant loss of Foxp3 protein expression in *Lkb1*-deficient T_{reg} cells after 4 days of culture (Fig. 4c). Strikingly, addition of IL-12, a potent STAT4 activator, caused the loss of

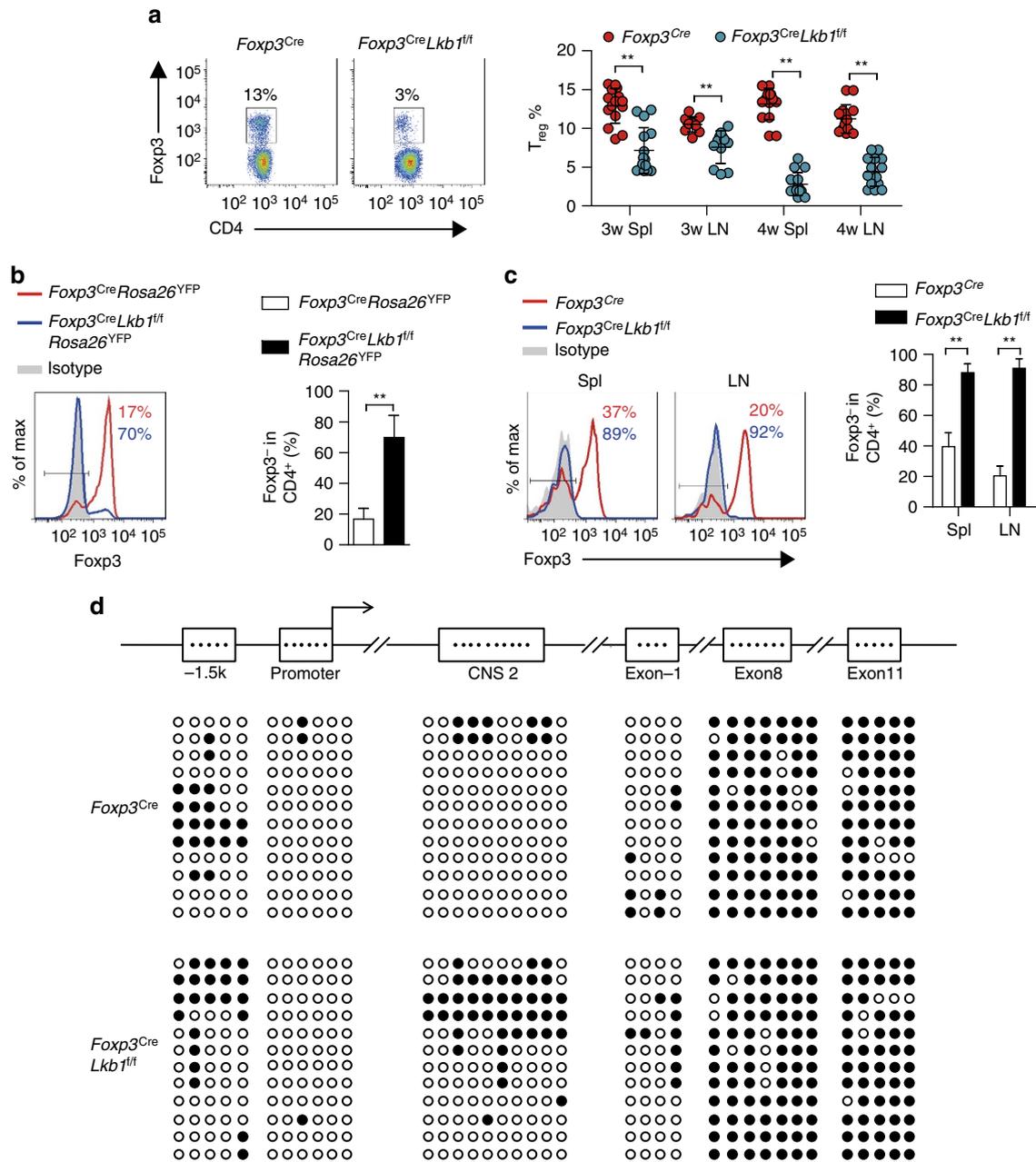


Figure 3 | Lkb1 stabilizes Foxp3 expression and prevents CNS2 methylation. (a) Foxp3⁺ cell percentages among CD4⁺ T cells in spleen and lymph nodes from *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice of different ages ($n = 10-15$). (b) Foxp3 expression in CD4⁺ *Rosa26^{YFP}* cells (*Rosa26^{YFP}* labelled the cells that have experienced the expression of Foxp3, *Rosa26^{YFP}* was much brighter than and thus distinguishable from Foxp3-YFP cells from *Foxp3^{Cre}Rosa26^{YFP}* and *Foxp3^{Cre}Lkb1^{fl/fl}Rosa26^{YFP}* mice). (c) Foxp3 expression in T_{reg} cells doubly sorted from *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice and transferred together with T_n cells into *Rag1^{-/-}* mice for 3 weeks ($n = 3$). (d) Methylation of the CpG motifs of the -1.5 kb region, promoter, CNS2, exon 1, exon 8 and exon 11 across the *Foxp3* locus in T_{reg} cells from *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice, determined by bisulfite sequencing. Filled circles represent methylated CpG sites and open circles represent unmethylated CpG sites. Two-way analysis of variance (ANOVA) was used for statistical analyses in **a,c**, and unpaired two-tailed Student's *t*-test was used for statistical analyses in **b** (** $P < 0.01$); error bars represent s.d.; all data are representative of at least two independent experiments.

Foxp3 expression in more than 50% Lkb1-deficient but not wild-type T_{reg} cells (Fig. 4c), suggesting that STAT4 hyperactivation is the causative reason for the instability of Lkb1-deficient T_{reg} cells. In addition, the apoptosis and proliferation were barely changed in Lkb1-deficient T_{reg} cells compared with wild-type T_{reg} cells (Supplementary Fig. 4c,d), suggesting that the loss of Foxp3 expressing Lkb1-deficient T_{reg} cells in response to IL-2 + IL-12 is not due to their altered apoptosis or proliferation. Intriguingly, both divided and nondivided Lkb1-deficient T_{reg} cells lost their stability (Fig. 4d). To determine whether the

instability of Lkb1-deficient T_{reg} cells was due to an intrinsic signalling alteration or *in vivo* selection of the T_{reg} subpopulations with reduced stability, we used *ERT2^{Cre}Lkb1^{fl/fl}Rosa26^{YFP}* mice in which Lkb1 could be induced to be acutely deleted in T_{reg} cells (Supplementary Figs 4e and 8l,m), thus avoiding the long-term *in vivo* selection. A similar IL-12-induced loss of stability was observed in T_{reg} cells after induced Lkb1 deletion with 4-hydroxytamoxifen *in vitro* (Fig. 4e), implying an intrinsic role for STAT4 hyperactivation in destabilizing Lkb1-deficient T_{reg} cells.

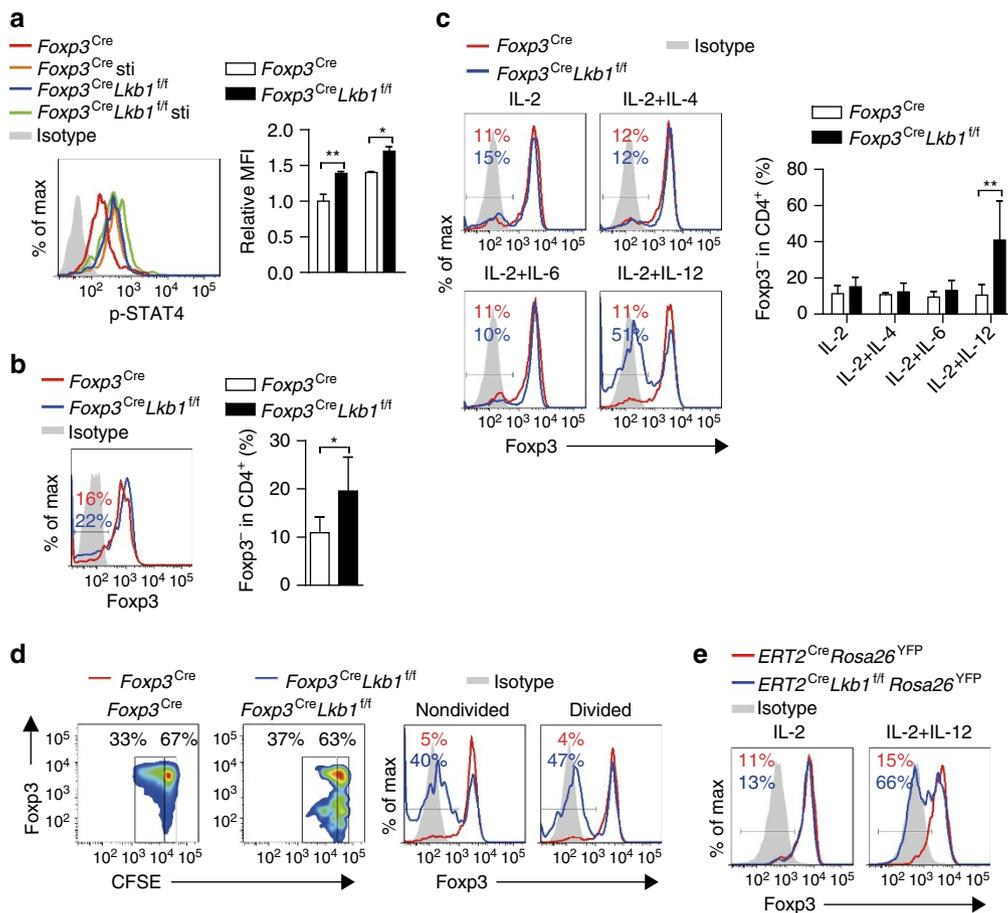


Figure 4 | Lkb1 prevents STAT4 activation to maintain T_{reg} cell stability. (a) Intracellular phosphorylated STAT4 in *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} T_{reg} cells treated with or without IL-12 ($n = 3$). (b) Foxp3 expression in *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} T_{reg} cells co-cultured with DCs without any cytokines ($n = 7$). (c) Foxp3 expression in *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} T_{reg} cells co-cultured with DCs supplemented with indicated cytokines ($n = 3$). (d) Foxp3 expression in divided and nondivided *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} T_{reg} cells co-cultured with DCs for 4 days supplemented with IL-2 and IL-12. (e) Foxp3 expression in *ERT2*^{Cre}*Rosa26*^{YFP} and *ERT2*^{Cre}*Lkb1*^{fl/fl}*Rosa26*^{YFP} T_{reg} cells co-cultured with DCs supplemented with indicated cytokines and 4-hydroxytamoxifen. Two-way analysis of variance (ANOVA) was used for statistical analyses in **a,c**, and unpaired two-tailed Student's *t*-test was used for statistical analyses in **b** (* $P < 0.05$, ** $P < 0.01$); error bars represent s.d.; all data are representative of at least two independent experiments.

We then tested whether STAT4 could bind to and induce methylation of the *Foxp3* locus in *Lkb1*-deficient T_{reg} cells. Chromatin immunoprecipitation (ChIP) experiments showed that *Lkb1*-deficient T_{reg} cells had significantly more STAT4 binding to the CNS2 than did the wild-type cells in response to IL-2 and IL-12 (Fig. 5a and Supplementary Datas 1 and 2). Additionally, compared with IL-2 treatment alone, IL-2 and IL-12 treatment decreased STAT5 binding to the CNS2 in *Lkb1*-deficient T_{reg} cells (Fig. 5a), suggesting that STAT4 might outcompete STAT5 in DNA binding in the *Foxp3* locus. Since DNA methyltransferase (Dnmt) 1 and 3a were critical for maintaining and inducing DNA methylation respectively³⁷, we tested whether STAT4 was capable of recruiting Dnmt1 or 3a to the *Foxp3* locus. *Lkb1*-deficient T_{reg} cells had significantly more Dnmt1 binding to the CNS2 than did the wild-type cells in response to IL-2 and IL-12 (Fig. 5b), consistent with the results of STAT4 (Fig. 5a). Notably, STAT4 associated with Dnmt1 but not Dnmt3a in T_{reg} cells (Fig. 5c and Supplementary Fig. 8n,o). Furthermore, we examined the methylation of CNS2 in *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} T_{reg} cells treated with IL-2, IL-2 + IL-12 or IL-2 + IL-12 + 5-aza-deoxycytidine (5-aza-dC, a DNA methyltransferase inhibitor) to investigate the relationship between STAT4 and CNS2 methylation. In *Foxp3*^{Cre}*Lkb1*^{fl/fl} T_{reg} cells, the methylation of CNS2 was higher after the treatment of

IL-2 + IL-12 compared with the treatment of IL-2 that could be reversed by 5-aza-dC (Fig. 5d). Indeed, the addition of 5-aza-dC rescued the loss of Foxp3 expression in *Lkb1*-deficient T_{reg} cells (Fig. 5e). Since 5-aza-dC changes the global DNA methylation status, leading to the global changes in gene expression, we cannot exclude its effect on the expression of other genes. Nevertheless, these results together suggest that *Lkb1* maintains stable Foxp3 expression in T_{reg} cells under inflammatory conditions by preventing STAT4-mediated recruitment of Dnmt1 to and subsequent methylation of the *Foxp3* CNS2.

Lkb1 restrains STAT4 and NF- κ B signalling. The *Stat4* and *Il12rb2* mRNA and IL-12R β 2 protein expression levels were elevated in *Lkb1*-deficient T_{reg} cells that could explain the increased STAT4 phosphorylation in these cells in response to IL-12 (Supplementary Fig. 5a,b). To determine which signalling promotes *Stat4* and *Il12rb2* expression in *Lkb1*-deficient T_{reg} cells, we examined the activation of NF- κ B and AKT, the major signalling mediators driving T helper type cell development^{38,39}. We found that NF- κ B p65 but not AKT was hyperactivated in *Lkb1*-deficient T_{reg} cells (Fig. 6a and Supplementary Figs 5c,d and 8p-r), and the chemical inhibitors to NF- κ B reduced *Stat4* and *Il12rb2* mRNA expression (Fig. 6b), as well as STAT4

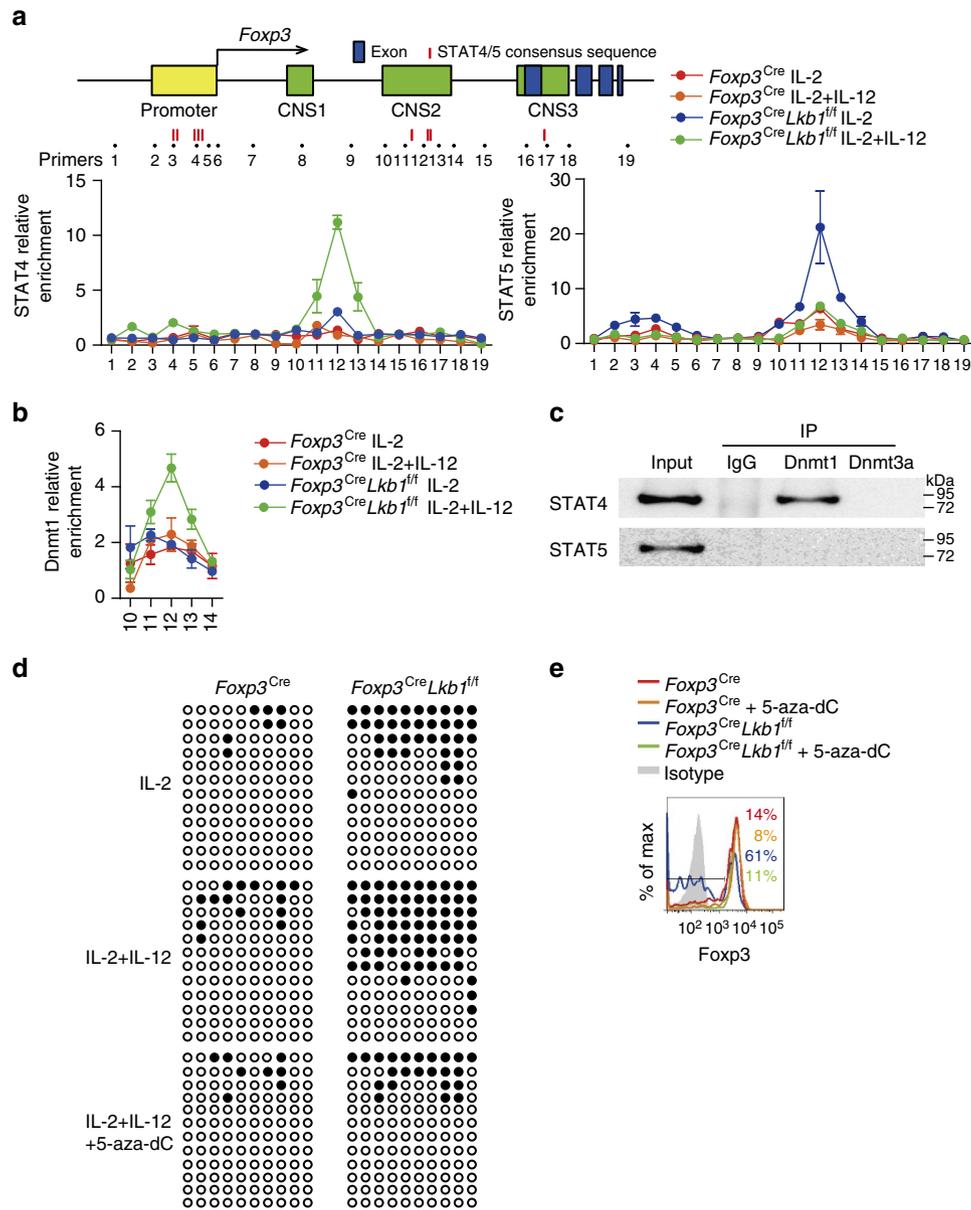


Figure 5 | Lkb1 prevents STAT4 binding to CNS2. (a) Binding of STAT4 and STAT5 to the *Foxp3* locus in *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} T_{reg} cells treated with IL-2 or IL-2 plus IL-12, determined by ChIP. (b) Binding of Dnmt1 to the CNS2 of *Foxp3* locus in *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} T_{reg} cells treated with IL-2 or IL-2 plus IL-12, determined by ChIP. (c) STAT4 and STAT5 coprecipitation with Dnmt1 and Dnmt3a was analysed by using nuclear extract from *in vitro* expanded T_{reg} cells. (d) Methylation of the CpG motifs of the *Foxp3* promoter and CNS2 in *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} T_{reg} cells determined by bisulfite sequencing, and T_{reg} cells were treated with indicated reagents. Filled circles represent methylated CpG sites and open circles represent unmethylated CpG sites. (e) *Foxp3* expression in *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} T_{reg} cells co-cultured with DCs supplemented with IL-2 plus IL-12 in the presence or absence of 5-aza-dC. All data are representative of at least two independent experiments.

phosphorylation (Fig. 6c). In addition, conserved NF-κB binding sites were present at the *Stat4* and *Il12rb2* loci (Supplementary Data 1), and ChIP experiments demonstrated more binding of NF-κB p65 to *Stat4* and *Il12rb2* in *Lkb1*-deficient T_{reg} cells than in wild-type cells (Fig. 6d and Supplementary Datas 1 and 2), suggesting a direct role of NF-κB activation in promoting *Stat4* and *Il12rb2* transcription. Of note, chemical inhibitors of NF-κB could partially rescue the instability of *Lkb1*-deficient T_{reg} cells (Fig. 6e). These results collectively indicate an involvement of NF-κB activation in driving STAT4 activation and instability of *Lkb1*-deficient T_{reg} cells.

IκB (inhibitor of NF-κB) kinase (IKK) phosphorylates IκB, leading to their degradation and activation of transcription factor NF-κB. IKK is composed of three subunits, the similar protein

kinases IKKα and IKKβ, and a regulatory subunit IKKγ⁴⁰. The phosphorylations serines (Ser) in IKKα (Ser 176/180)/β (Ser 177/181) and IκBα (Ser 32/36) were all increased in *Lkb1*-deficient T_{reg} cells (Fig. 6f,g), suggesting that *Lkb1* might dampen NF-κB signalling by inhibiting IKKα/β activation.

Lkb1 promotes the expression of immunosuppressive genes.

We further explored whether there are defects in *Lkb1*-deficient T_{reg} cells that did not lose *Foxp3* expression yet. The T-cell activation phenotype was already obvious (Fig. 7a,b) in 9–11-day-old *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice that had normal percentages of T_{reg} cells among CD4⁺ T cells (Fig. 7c), suggesting that the suppressor function of *Lkb1*-deficient T_{reg} cells might be impaired.

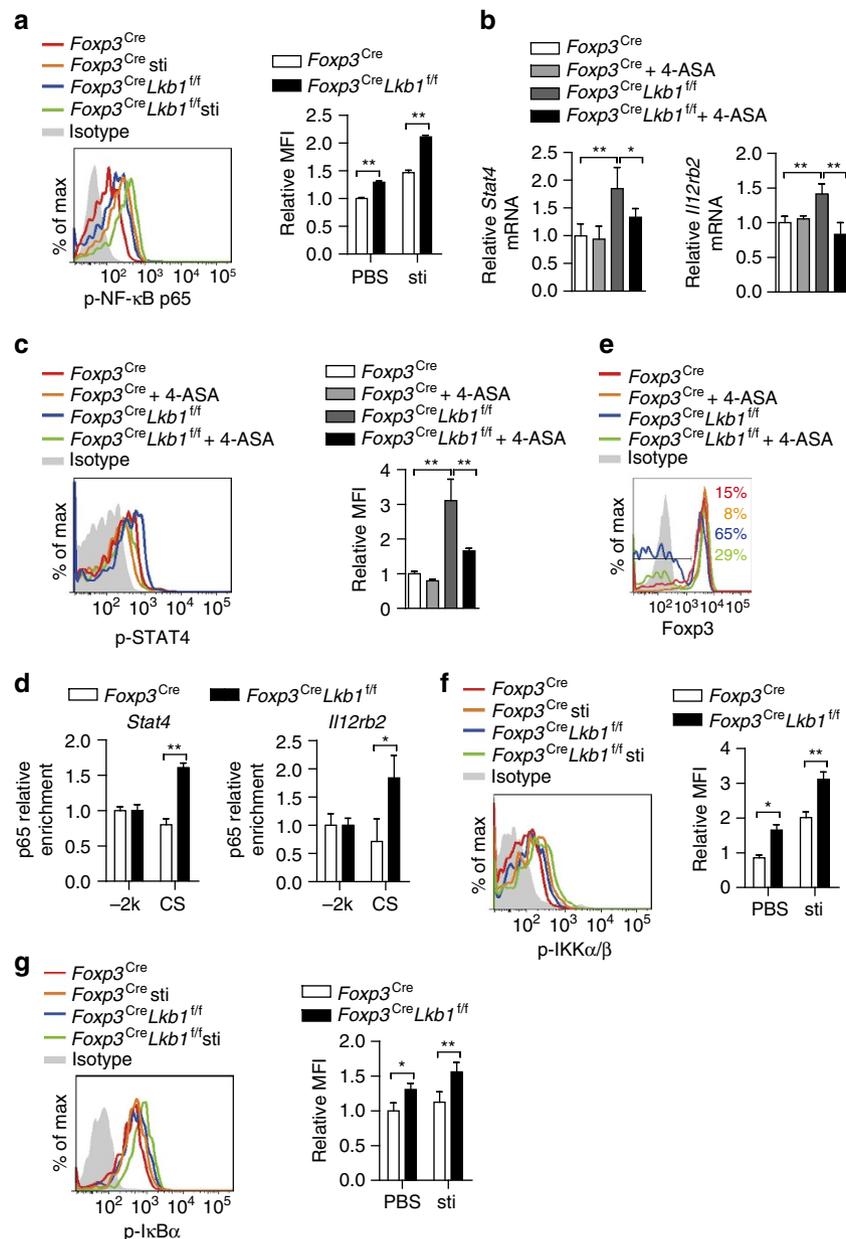


Figure 6 | Lkb1 suppresses NF-κB signaling to restrain STAT4 activation. (a) Intracellular phosphorylated NF-κB p65 in *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* T_{reg} cells with or without IL-2 stimulation ($n = 3$). (b) *Stat4* and *Il12rb2* mRNA in *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* T_{reg} cells supplemented with or without 4-ASA for 16 h ($n = 3$). (c) Intracellular phosphorylated STAT4 in *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* T_{reg} cells supplemented with or without 4-ASA. (d) P65 enrichment to consensus sequences (CS) in *Stat4* and *Il12rb2* locus in *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* T_{reg} cells determined by ChIP. (e) Foxp3 expression in *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* T_{reg} cells co-cultured with DCs supplemented with IL-2 and IL-12, with or without 4-ASA. (f) Intracellular phosphorylated IKKα (Ser176/180)/β (Ser177/181) in *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* T_{reg} cells with or without IL-2 stimulation ($n = 3$). (g) Intracellular phosphorylated IκBα (Ser 32/36) in *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* T_{reg} cells with or without IL-2 stimulation ($n = 3$). Two-way analysis of variance (ANOVA) was used for statistical analyses in (a,b,c,d,f,g) (* $P < 0.05$, ** $P < 0.01$); error bars represent s.d.; all data are representative of at least two independent experiments.

Indeed, *in vitro* T_{reg} suppression assay showed that with different T_{reg} to T_n ratios, Lkb1-deficient T_{reg} cells could not efficiently suppress the proliferation of responder T cells (Fig. 7d), indicating an impaired suppressive capacity of Lkb1-deficient T_{reg} cells.

To investigate the molecular mechanisms by which Lkb1 controls T_{reg} cell function, we compared the transcriptional profiling of $CD4^+ YFP^+$ T_{reg} cells sorted from *Foxp3^{Cre/+}* mice with that from *Foxp3^{Cre/+}Lkb1^{fl/fl}* mice lacking autoimmune diseases, thus avoiding the secondary impact of the inflammatory environment on T_{reg} cell gene expression (Supplementary Data 3). Kyoto Encyclopedia of Genes and Genomes (KEGG)

pathway analysis revealed that the most altered pathways were enriched for genes regulating immune function (Supplementary Fig. 6a). Intriguingly, Lkb1-deficient T_{reg} cells have reduced expression of a wide variety of genes critically involved in T_{reg} cell suppressor function including those encoding secreted molecules associated with immune suppression (*Fgl2*, *Il10* and *Il18*)^{11,41}, factors in wound-healing processes (*Tfpi*)¹¹, regulators of reactive oxygen generation (*Cybb*)¹¹, enzymes catalysing the generation of adenosine that directly inhibit proliferation of effector T cells (*Entpd1* and *Nt5e*)^{11,42}, chemokine receptors critical for T_{reg} cell migration (*Ccr6*)⁴³ and cell surface molecules facilitating

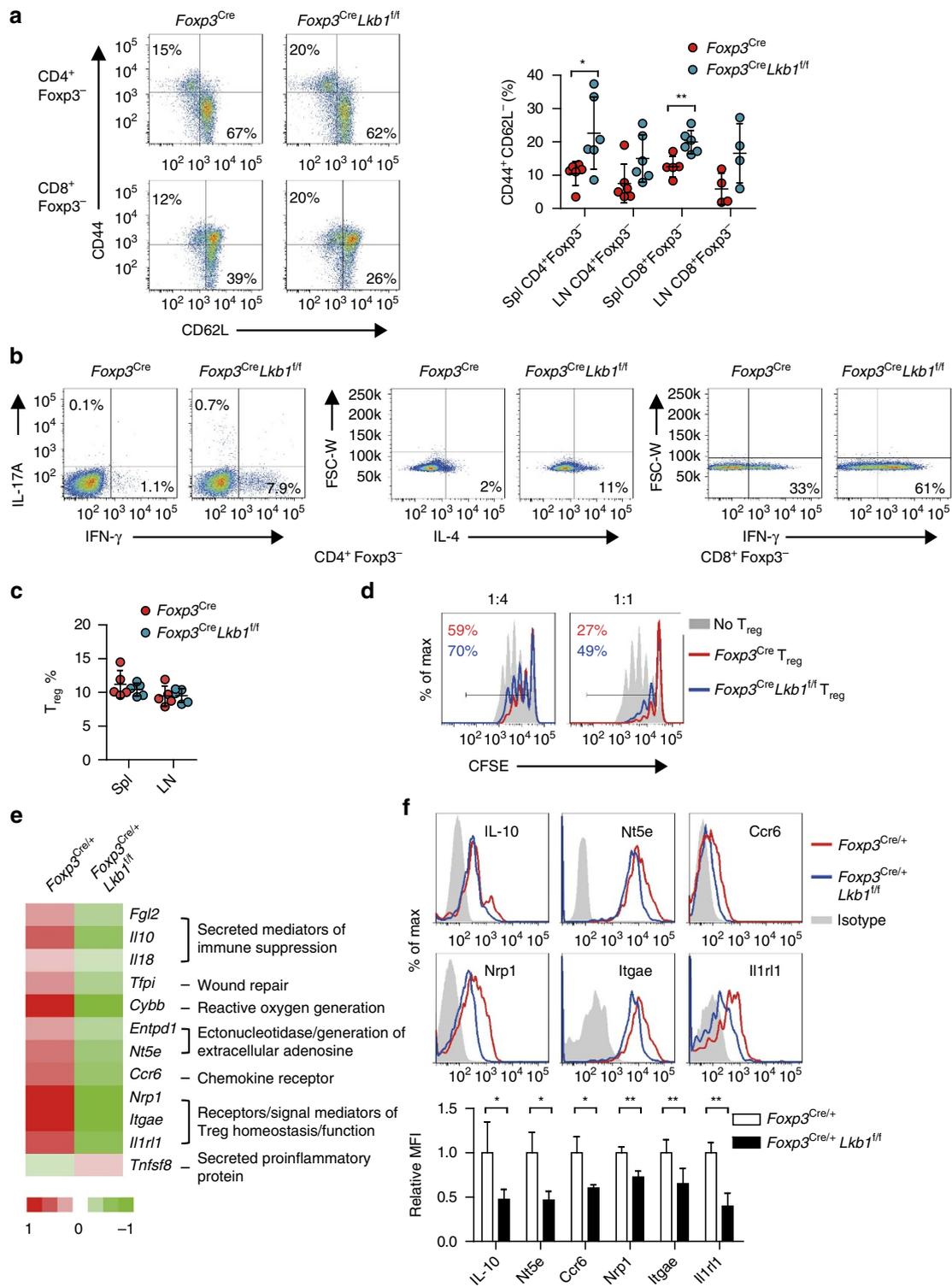


Figure 7 | *Lkb1* promotes the expression of diverse immunosuppressive genes. (a) CD44^{high}CD62L^{low} effector/memory cells among splenic CD4⁺Foxp3⁻ and CD8⁺Foxp3⁻ T cells from 9–11-day-old *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice (n = 4–6). (b) Intracellular staining of cytokines in splenic CD4⁺Foxp3⁻ and CD8⁺Foxp3⁻ T cells from 9–11-day-old *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice stimulated with phorbol myristate acetate (PMA) and ionomycin for 4 h. (c) Percentages of Foxp3⁺ T_{reg} cells among CD4⁺ T cells in the spleen and lymph nodes from 9–11-day-old *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice (n = 5). (d) Suppression of proliferation of CFSE-labelled T_n cells (responding cells, T_{resp}) by different ratios of CD4⁺YFP⁺ T_{reg} cells from *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice. T_{resp} cell division was determined by CFSE dilution at the indicated ratios of cell numbers between T_{reg} cell and T_{resp} cells. The experiment was repeated three times. (e) T_{reg} cell function-related genes differentially expressed between CD4⁺YFP⁺ T_{reg} cells from *Foxp3*^{Cre/+} and *Foxp3*^{Cre/+}*Lkb1*^{fl/fl} mice determined by transcriptional profiling, shown in groups based on their functions. Fold difference means fold change of gene expression levels in T_{reg} cells from *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice compared with that from *Foxp3*^{Cre} mice. (f) Expression of indicated proteins on splenic T_{reg} cells from *Foxp3*^{Cre/+} and *Foxp3*^{Cre/+}*Lkb1*^{fl/fl} mice (n = 3). Two-way analysis of variance (ANOVA) was used for statistical analyses in (a, c, f) (*P < 0.05, **P < 0.01); error bars represent s.d.; all data are representative of at least two independent experiments.

T_{reg} cell suppressor functions in either tumour environment (*Nrp1*)⁴⁴ or organ-specific inflammatory conditions (*Itgae* and *Il1rl1*)^{45,46} (Fig. 7e and Supplementary Fig. 6b). On the other hand, the loss of *Lkb1* promoted the expression of genes encoding proinflammatory mediators (*Tnfsf8*)¹¹ (Fig. 7e and Supplementary Fig. 6b). It was further confirmed that *Lkb1*-deficient T_{reg} cells from *Foxp3*^{Cre/+}*Lkb1*^{fl/fl} mice had decreased protein levels of IL-10, Nt5e, Ccr6, Nrp1, Itgae and Il1rl1 (Fig. 7f). Most gene expression alterations were also recapitulated in T_{reg} cells from *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice (Supplementary Fig. 6b). These results clearly showed that *Lkb1* promoted the immunosuppressive programs in T_{reg} cells.

TGF- β signalling in *Lkb1*-implemented T_{reg} cell function.

TGF- β signalling plays crucial roles in T_{reg} cell generation and function⁴⁷. Transcriptional profiling analysis showed that *Lkb1*-deficient T_{reg} cells express lower levels of *Tgfb1* and *Tgfb2* mRNA that was confirmed by real-time PCR and western blot (Fig. 8a and Supplementary Figs 7a and 8s,t). Hence, we proposed that this pathway might be defective in *Lkb1*-deficient T_{reg} cells. Indeed, Smad2 (Ser 456/457)/Smad3 (Ser 423/425) phosphorylation was decreased (Fig. 8b), and Smad-DNA binding assay indicated that Smad transcriptional activity was impaired in *Lkb1*-deficient T_{reg} cells (Fig. 8c). To determine whether impaired TGF- β signalling contributed to the impairment of *Lkb1*-deficient T_{reg} cells, we conducted loss-of-function experiments by generating a mouse line with TGF- β 2 specifically deleted in T_{reg} cells (*Foxp3*^{Cre}*Tgfb2*^{fl/fl}, Supplementary Figs 7b and 8u,v) that displayed decreased Smad activation (Supplementary Fig. 7c). Although *Foxp3*^{Cre}*Tgfb2*^{fl/fl} mice did not show overt autoimmune disease, increased percentages of CD44^{high}CD62L^{low} effector/memory T cells (Fig. 8d) were observed despite a slightly increased T_{reg} cell frequency among CD4⁺ T cells (Fig. 8e), indicating the functional impairment of TGF- β 2-deficient T_{reg} cells. Consistently, TGF- β 2-deficient T_{reg} cells displayed impaired suppressive capacity *in vitro* (Fig. 8f). To determine the global effects of defective TGF- β signalling on gene transcription, we conducted transcriptional profiling and found that TGF- β 2 deficiency led to the decreased expression of many genes important for T_{reg} cell suppressor function (Supplementary Data 4). Remarkably, TGF- β 2 deficiency recapitulated part of the suppressor gene expression alterations in *Lkb1*-deficient T_{reg} cells, including *Nt5e*, *Ccr6*, *Nrp1*, *Itgae* and *Il1rl1* (Fig. 8g and Supplementary Data 4), that was also confirmed at protein levels (Fig. 8h,i). However, the *Lkb1*-regulated *Fgl2*, *Il10*, *Tfpi*, *Entpd1* and *Tnfsf8* transcripts were not altered in TGF- β 2-deficient T_{reg} cells, suggesting that *Lkb1* also utilize TGF- β -independent mechanisms to promote T_{reg} cell function. To further determine whether lower expression of TGF- β 2 was the causative reason for impaired suppression function of *Lkb1*-deficient T_{reg} cells, we generated T_{reg} cells (from *Foxp3*^{Cre} and *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice) transduced with retrovirus carrying TGF- β 2 complementary DNA (cDNA). TGF- β 2 was successfully expressed in RFP⁺YFP⁺ T_{reg} cells (Supplementary Fig. 7c). Overexpression of TGF- β 2 could partially rescue the expression of suppressor genes and suppression function of *Lkb1*-deficient T_{reg} cells (Fig. 8j and Supplementary Fig. 7d,e). Together, these results indicate that *Lkb1* implements T_{reg} cell function partially through promoting TGF- β signalling. TGF- β 2-deficient T_{reg} cells did not lose stability when co-transferred with T_n cells into *Rag1*^{-/-} mice (Fig. 8k), suggesting that the decreased expression of TGF- β receptors does not account for the instability of *Lkb1*-deficient T_{reg} cells.

Discussion

Here, we demonstrate that mice with T_{reg} cell-specific deletion of *Lkb1* develop a fatal, early-onset, autoimmune disease with moribund at ~35 days of age. Such a fulminant disease is comparable to that observed in mice devoid of T_{reg} cells or deficient in *Foxp3*. Similar devastating diseases have also been reported to be present in mice with selective deficiency of Foxo1 (ref. 48) or Raptor, the rapamycin sensitive partner of mTORC1 (mammalian target of rapamycin complex 1)²⁶ in T_{reg} cells, due to impaired T_{reg} cell function. Intriguingly, *Lkb1*-deficient T_{reg} cells lost their lineage-specific features and exhibited severe impairment in the maintenance of *Foxp3* expression and suppressor function that may underlie the extremely severe autoimmunity in *Foxp3*^{Cre}*Lkb1*^{fl/fl} mice. Although *Lkb1* expression is not restricted to T_{reg} cells, T_{reg} cells substantially upregulate *Lkb1* protein expression upon TCR stimulation. Given that TCR signalling is essential for T_{reg} cell homeostasis and function, TCR-induced upregulation of *Lkb1* protein may represent a critical feed-forward loop to stabilize T_{reg} cell lineage identity.

It is generally recognized that *Foxp3* expression is highly stable in T_{reg} cells. Whether a small proportion of T_{reg} cells can lose *Foxp3* expression in certain conditions is a controversial topic⁶. Some studies showed that T_{reg} cells could lose *Foxp3* expression and become 'ex- T_{reg} ' cells under certain conditions²⁵, while others argued that the so-called 'ex- T_{reg} ' cells just might be the conventional T cells that transiently express *Foxp3*, namely poorly committed T_{reg} cells²⁷. Here, we showed that deficiency of *Lkb1* caused very severe defects in T_{reg} cell stability. More than 70% of T_{reg} cells lost stability in *Foxp3*^{Cre}*Lkb1*^{fl/fl}*Rosa26*^{YFP} mice. In addition, the majority of the *Lkb1*-deficient T_{reg} cells lost *Foxp3* expression after being transferred into *Rag1*^{-/-} mice, indicating that *Lkb1*-deficient mature T_{reg} cells continuously lost *Foxp3* expression, rather than that a subset of *Lkb1*-deficient T cells only transiently expressed *Foxp3* (poorly committed T_{reg} cells). Such a severe defect was rarely reported in literatures, placing *Lkb1* as a particularly important maintainer of *Foxp3* stability. The demethylated status of *Foxp3* CNS2 is critically involved in the maintenance of stable *Foxp3* expression in T_{reg} cells^{8-10,28-32}. Substantial methylation of CNS2 but not promoter was observed in *Lkb1*-deficient T_{reg} cells. Treatment of IL-12 further enhanced the methylation of CNS2 and caused the loss of *Foxp3* expression in *Lkb1*-deficient but not wild-type T_{reg} cells. Moreover, the DNA methyltransferase inhibitor 5-aza-dC reduced the methylation of CNS2 and rescued the instability of *Lkb1*-deficient T_{reg} cells. These results suggest that *Lkb1* stabilizes *Foxp3* expression by building up the demethylated status of CNS2. It has been recently reported that deletion of CNS2 caused cell-division-dependent loss of *Foxp3* expression^{8,10}. In contrast, both divided and nondivided *Lkb1*-deficient T_{reg} cells lost *Foxp3* expression, suggesting that *Lkb1* might also promote *Foxp3* expression by CNS2-independent mechanisms, which need future exploration.

The role of STAT4 in T_{reg} cells is not well defined. Here we showed that deficiency of *Lkb1* in T_{reg} cells resulted in markedly more IL-12-induced STAT4 activation and STAT4 binding to CNS2. STAT4 associated with Dnmt1 and DNA methyltransferase chemical inhibitor rescued the instability of *Lkb1*-deficient T_{reg} cells in response to IL-12. Thus, *Lkb1* suppresses IL-12/STAT4 activation to stabilize *Foxp3* expression in T_{reg} cells. We further showed that NF- κ B signalling is activated in *Lkb1*-deficient T_{reg} cells, that increased *Il12rb2* and *Stat4* mRNA expression and STAT4 activation. However, inhibition of NF- κ B signalling can only partially rescue the instability of *Lkb1*-deficient T_{reg} cells, implicating that NF- κ B-independent signalling mechanisms might also contribute to the unstable *Foxp3* expression in *Lkb1*-deficient T_{reg} cells.

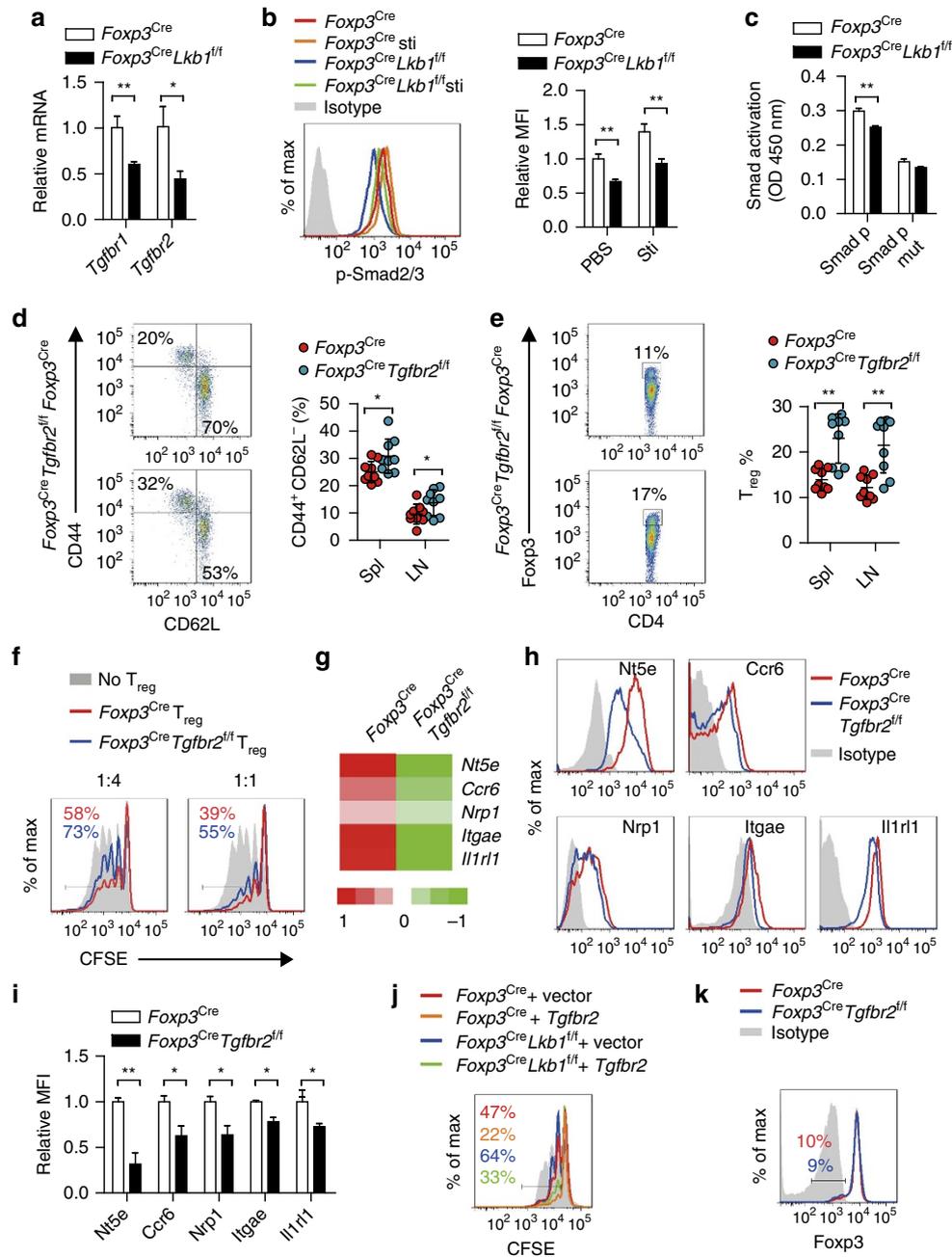


Figure 8 | TGF- β signaling is involved in Lkb1-implemented T_{reg} cell suppressor function. (a) *Tgfb1* and *Tgfb2* mRNA expression in *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* T_{reg} cells by real-time PCR ($n = 3$). (b) Intracellular expression of phosphorylated Smad2/3 in T_{reg} cells from *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice, with or without TGF- β stimulation ($n = 3$). (c) Binding capacity of Smad2/3 to recognized DNA sequences in nuclear lysates extracted from T_{reg} cells from *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice. (d) CD44 and CD62L expression on CD4⁺Fopx3⁻ T cells from 6-week-old *Foxp3^{Cre}* and *Foxp3^{Cre}Tgfb2^{fl/fl}* mice ($n = 6-11$). (e) Fopx3 expression in CD4⁺ T cells from 6-week-old *Foxp3^{Cre}* and *Foxp3^{Cre}Tgfb2^{fl/fl}* mice ($n = 6-11$). (f) Suppression of proliferation of T_{reg} cells labelled with CFSE by different ratios of CD4⁺YFP⁺ T_{reg} cells from *Foxp3^{Cre}* and *Foxp3^{Cre}Tgfb2^{fl/fl}* mice. The experiment was repeated three times. (g) Some genes differentially expressed between CD4⁺YFP⁺ T_{reg} cells from *Foxp3^{Cre}* and *Foxp3^{Cre}Tgfb2^{fl/fl}* T_{reg} cells determined by transcriptional profiling. Fold difference means fold change of gene expression levels in T_{reg} cells from *Foxp3^{Cre}Tgfb2^{fl/fl}* mice compared with that from *Foxp3^{Cre}* mice. (h,i) Expression of indicated proteins on splenic T_{reg} cells from *Foxp3^{Cre}* and *Foxp3^{Cre}Tgfb2^{fl/fl}* mice ($n = 3$). (j) Suppression of proliferation of T_{reg} cells labelled with CFSE by different ratios of CD4⁺YFP⁺ T_{reg} cells from *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice that were transduced with retrovirus carrying TGF- β 2 or control vector. (k) Fopx3 expression in CD4⁺YFP⁺ T_{reg} cells doubly sorted from *Foxp3^{Cre}* and *Foxp3^{Cre}Tgfb2^{fl/fl}* mice and transferred together with T_{reg} cells into *Rag1^{-/-}* mice for 3 weeks. Two-way analysis of variance (ANOVA) was used for statistical analyses in (a,b,c,d,e,i) (* $P < 0.05$, ** $P < 0.01$); error bars represent s.d.; all data are representative of at least two independent experiments.

Independent of maintaining Fopx3 expression, Lkb1 also promotes T_{reg} cell-suppressive capacity. Loss of Lkb1 in T_{reg} cells results in the moderate downregulation of a number of immunosuppressive genes (*Fgl2*, *Tfpi*, *Cybb*, *Entpd1*, *Nt5e*, *Nrp1*,

Itgae, *Il18*, *Il10*, *Ccr6*, *Il1r1*), suggesting that Lkb1 implements T_{reg} cell function through broadly promoting T_{reg} suppressor gene expression program. Lkb1 has been previously shown to positively or negatively regulate TGF- β /Smad signalling in a cell

type-dependent manner^{49,50}, but the precise mechanism is not fully elucidated. Here, we found the effect of *Lkb1* on promoting T_{reg} suppressor function was partially dependent on its augmentation of TGF- β signalling. TGF- β pathway is important for T_{reg} cell development in the thymus and periphery^{47,51,52}, but whether and how it affects T_{reg} cell function remain poorly understood. Our results demonstrate that TGF- β 2-deficient T_{reg} cells have impaired suppressive capacity associated with decreased expression of a number of genes associated with immune suppression (*Nt5e*, *Ccr6*, *Nrp1*, *Itgae* and *Il1rl1*), partially recapturing the suppressor gene expression alterations in *Lkb1*-deficient T_{reg} cells. Furthermore, overexpression of TGF- β 2 partially rescued the expression of certain suppressor genes and the suppressor function of *Lkb1*-deficient T_{reg} cells, supporting that impaired TGF- β signalling contributed to the impaired suppressor function of *Lkb1*-deficient T_{reg} cells. However, numerous *Lkb1*-regulated genes, including *Fgl2*, *Tfpi*, *Cybb*, *Entpd1* and *Il18*, were not changed in TGF- β 2-deficient T_{reg} cells, suggesting that TGF- β -independent mechanisms were also applied by *Lkb1* to promote T_{reg} cell suppressor function.

AMPK-induced fatty acid oxidation is a critical metabolic feature of T_{reg} cells⁵³. To our surprise, AMPK activation is not dramatically altered in *Lkb1*-deficient T_{reg} cells, and T_{reg} cell-specific deletion of both AMPK1 α and AMPK2 α does not cause T_{reg} cell abnormality and immune disturbance, suggesting that *Lkb1* does not control T_{reg} cells by promoting AMPK activity in steady-state conditions.

Lkb1 has been previously shown to promote thymocytes development, maintain survival and proliferation of peripheral T cells and restrain peripheral conventional T-cell activation and proinflammatory cytokine production^{15,16,54} (we got similar results in certain experiments after inducing *Lkb1* deletion in conventional T cells from *ERT2^{Cre}Lkb1^{fl/fl}* mice, Supplementary Fig. 7f,g). Here, we find that *Lkb1* maintains the lineage identity of T_{reg} cells through mechanisms involving the regulation of NF- κ B/STAT4 and TGF- β signalling (Fig. 8). NF- κ B/STAT4 and TGF- β signalling are separately controlled by *Lkb1* (Supplementary Fig. 7h-j), suggesting a function of *Lkb1* in setting up the optimal activation thresholds of multiple intracellular signalling in T_{reg} cells. Furthermore, the upregulation of *Lkb1* protein by TCR signalling in T_{reg} but not in conventional T cells indicates a specific regulation of *Lkb1* expression required for preserving the T_{reg} cell lineage. Future studies will determine whether and how the environmental cues change *Lkb1* in T_{reg} cells to affect their stability/function under physiological and pathological conditions, and whether *Lkb1* can be targeted to treat T_{reg} -cell-related immune diseases.

Methods

Mice. All animals were maintained in specific pathogen-free barrier facilities and were used in accordance with protocols approved by the institutional animal care and user committee at the Institute of Hematology, Chinese Academy of Medical Sciences. C57BL/6, CD45.1⁺, *Lkb1^{fl/fl}*, *AMPK α 1^{fl/fl}*, *AMPK α 2^{fl/fl}*, *Tgfb2^{fl/fl}*, *ERT2^{Cre}*, *Foxp3^{YFP-Cre}* (*Foxp3^{Cre}*), *Rosa26^{YFP}* and *Rag1^{-/-}* mice were purchased from Jackson Laboratories. All mice had been backcrossed with C57BL/6 mice for at least 7 generations. *Lkb1^{fl/fl}*, *AMPK α 1^{fl/fl}*, *AMPK α 2^{fl/fl}*, *Tgfb2^{fl/fl}* mice were crossed with *Foxp3^{Cre}* mice to generate *Foxp3^{Cre}Lkb1^{fl/fl}*, *Foxp3^{Cre}AMPK α 1^{fl/fl}*, *Foxp3^{Cre}AMPK α 2^{fl/fl}* and *Foxp3^{Cre}Tgfb2^{fl/fl}* mice, respectively. *Foxp3^{Cre}+Lkb1^{fl/fl}* mice were crossed with *Rosa26^{YFP}* mice to generate *Foxp3^{Cre}Lkb1^{fl/fl}Rosa26^{YFP}* mice. *Lkb1^{fl/fl}* mice were crossed with *ERT2^{Cre}* and *Rosa26^{YFP}* mice to generate *ERT2^{Cre}Lkb1^{fl/fl}Rosa26^{YFP}* mice. Sample size for various animal experiments was chosen based on prior data generated in the laboratory, and no mice were excluded from experiments. The details of mice age and sex are provided in the figure legends.

Cell purification and flow cytometry. Single-cell suspensions were prepared from spleen and peripheral lymph nodes for staining or cell purification. CD4⁺ T cells and CD11c⁺ DCs were purified with Dynabeads Untouched Mouse CD4 Cells Kits (Invitrogen, 11415D) or CD11c MicroBeads (Miltenyi Biotec, 130-108-338),

respectively. Indicated T-cell populations were sorted from purified CD4⁺ T cells with FACSAria III (BD Biosciences), and the sorted populations were >98% pure unless otherwise specified. Flow cytometry of cell surface molecules was performed as previously described⁵⁵, and the antibody (Supplementary Data 5) was diluted as the instructions of the manufacturers suggested. Intracellular staining of Foxp3, cytokines (spleen cells were stimulated with phorbol myristate acetate (50 ng ml⁻¹) and ionomycin (500 ng ml⁻¹) for 4 h before analysis of cytokine expression in indicated populations) and other proteins were performed with Foxp3 staining kits (eBioscience, 00-5523). Intracellular staining of phosphorylated proteins was performed on cells fixed with methanol and permeabilized with Triton X-100. The antibodies were obtained from eBioscience, Biogen, BD Biosciences, Cell Signalling Technology, R&D Systems and Invitrogen, and listed in Supplementary Data 5. Flow cytometry data were acquired on LSR II, LSRFortessa (BD Biosciences) or FACSCanto II (BD Biosciences) and analysed with Flowjo software (Tree Star).

Cell culture. Sorted CD4⁺ YFP⁺ T_{reg} cells from *Foxp3^{Cre}Lkb1^{fl/fl}* and *Foxp3^{Cre}* mice were labelled with carboxyfluorescein succinimidyl ester (CFSE) Cell Proliferation Kits (Invitrogen, C34554). CFSE-labelled T_{reg} cells were co-cultured with CD11c⁺ cells purified from CD45.1⁺ mice, supplemented with indicated combinations of cytokines of recombinant murine IL-2, IL-12, IL-6 and IL-4 (all 100 ng ml⁻¹, PeproTech), NF- κ B inhibitors sodium 4-aminosalicylate (4-ASA) or pyrrolidinedithiocarbamic acid (PDT) or DNA methyltransferase inhibitor 5-aza-dC in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum (Gibco) for 4 days. For inducible *Lkb1* deletion, T_{reg} cells were respectively sorted from *ERT2^{Cre}Lkb1^{fl/fl}Rosa26^{YFP}* and *ERT2^{Cre}Rosa26^{YFP}* mice and cultured for 5 days as described above, with the addition of 4-hydroxytamoxifen (1 μ M, Sigma) in the culture. In other experiments, sorted T_{reg} cells were incubated in plates pre-coated with anti-CD3 (2 μ g ml⁻¹; 145-2C11; eBioscience) and anti-CD28 (2 μ g ml⁻¹; 45.21; eBioscience) in the presence of IL-2 (100 ng ml⁻¹) for the indicated days. CFSE profiles and Foxp3 expressions were examined by flow cytometry after culturing.

In vivo T_{reg} cell maintenance. CD4⁺ CD25⁺ YFP⁺ T_{reg} cells were doubly sorted (>99.5% pure) from 3-week-old CD45.1⁻ CD45.2⁺ *Foxp3^{Cre}Lkb1^{fl/fl}* and CD45.1⁺ CD45.2⁻ *Foxp3^{Cre}* mice, and 2 \times 10⁵ of each type of cells were mixed with 4 \times 10⁵ CD4⁺ CD25⁻ CD44^{low} CD62L^{high} T_n cells sorted from CD45.1⁺ CD45.2⁻ mice and transferred into the *Rag1^{-/-}* mice by intraperitoneal injection. The recipient mice were analysed 3 weeks after transfer.

In vitro T_{reg} cell suppression. CD4⁺ CD25⁻ CD44^{low} CD62L^{high} T_n cells sorted from CD45.1⁺ mice were labelled with CFSE and used as responder cells (T_{resp}). T_{resp} cells (5 \times 10⁴) were cultured for 72 h with DCs (1 \times 10⁵) and soluble anti-CD3 (2 μ g ml⁻¹) in the presence or absence of the indicated numbers of CD4⁺ YFP⁺ T_{reg} cells sorted from *Foxp3^{Cre}Lkb1^{fl/fl}* or *Foxp3^{Cre}* mice.

Foxp3 gene methylation assay. Genomic DNA from sorted cells was bisulfite converted with EZ DNA Methylation-Direct kits according to the manufacturer's protocol (Epigentek, P-1026-050). Methylation-specific PCR primers (listed in Supplementary Data 2) were used for amplification of the promoter and intron 1 of *Foxp3* (corresponding to *Foxp3* conserved noncoding sequence 2). PCR products were subcloned into pGEM-T Easy vectors (Promega) and sequenced.

Western blot. CD4⁺ YFP⁺ T_{reg} cells and CD4⁺ YFP⁻ conventional T cells were sorted from *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice and treated as indicated, and then cells were lysed and western blot was carried out as previously described⁵⁵ with antibodies to *Lkb1* (D60C5, Cell Signalling Technology), phosphorylated AMPK (40H9, Cell Signalling Technology), IKK β (2C8, Cell Signalling Technology), phosphorylated IKK α / β (16A6, Cell Signalling Technology), phosphorylated I κ B α (14D4, Cell Signalling Technology), phospho-NF- κ B p65 (93H1, Cell Signalling Technology), NF- κ B p65 (D14E12, Cell Signalling Technology), STAT4 (C46B10, Cell Signalling Technology), AMPK α (D63G4, Cell Signalling Technology), phospho-Acetyl-CoA Carboxylase (D7D11, Cell Signalling Technology), phospho-STAT4 (Abcam), TGF- β R2 (Abcam) and GAPDH (D16H11, Cell Signalling Technology).

Microarray and quantitative real-time PCR. CD4⁺ YFP⁺ T_{reg} cells were sorted from spleen and lymph nodes of mice for RNA extraction with Trizol reagent (Invitrogen). Total RNA was reverse transcribed, amplified, labelled and hybridized to Mouse Genome 2.0 arrays (Affymetrix). Microarray data sets were analysed with Agilent Genespring GS 11 software. RNA of different samples was obtained in the same manner as in microarray analysis, and real-time PCR was performed with SYBR Green PCR Master Mix (ABI) as previously described⁵⁵. The sequences of the primer pairs used are listed in Supplementary Data 2.

DNA binding ELISA. TransAM Flexi kits (Active Motif, 40098) were used to test the activity of Smad transcription factors according to the manufacturer's protocol.

Briefly, nuclear extract from stimulated T_{reg} cells sorted from *Foxp3^{Cre}* and *Foxp3^{Cre}Lkb1^{fl/fl}* mice were mixed with biotinylated oligos (listed in Supplementary Data 2) recognized by transcription factors of interest, and incubated in a streptavidin-coated plate. After washing, primary antibodies specific for the bound transcription factor and horseradish peroxidase-conjugated secondary antibodies were subsequently added, and transcription factor binding activities were quantified using a microplate reader (Synergy H4, BioTek).

Co-immunoprecipitation. Sorted wild-type CD4⁺YFP⁺ T_{reg} cells were expanded in plates coated with anti-CD3 (2 µg ml⁻¹) and anti-CD28 (2 µg ml⁻¹) in the presence of IL-2 (100 ng ml⁻¹) for 5 days and collected. Cell lysis was prepared using Co-immunoprecipitation kits according to the manufacturer's protocol (Sigma, FLAGITP1), and incubated with antibodies against Dnmt1 (60B1220.1, Abcam), Dnmt3a (Abcam) and protein G agarose (Cell Signalling Technology). Immunoprecipitated proteins were detected as described above with antibodies against STAT4 (C46B10, Cell Signalling Technology) and STAT5 (Cell Signalling Technology).

Chromatin immunoprecipitation. ChIP assays were performed using the ChIP kits (Active Motif, 53040) according to the manufacturer's protocol. Precipitated DNA and input DNA were assessed by real-time PCR using primers listed in Supplementary Data 2.

Retroviral transduction. TGF-βR2 cDNA was subcloned into the retroviral vector pMYs-IRES-RFP. Retroviruses were produced by transfection of the Plat-E cells with polyethylenimine. T_{reg} cells (YFP⁺) were cultured in plates coated with anti-CD3 and anti-CD28 transduced in virus-containing media supplemented with polybrene by centrifuging for 1 h at 500 RCF. After 24 h, cells were expanded and then RFP⁺ YFP⁺ cells were sorted before further analysis. The Plat-E cell line was provided by the Cell Resource Center, Institute of Hematology and Hospital of Blood Diseases, Chinese Academy of Medical Sciences/Peking Union Medical College. The Cell Resource Center confirmed the species origin with PCR and checked free of mycoplasma contamination by PCR and culture.

Histopathology. Skins, lungs, livers and stomachs were removed from 3–4-week-old mice. Samples were formalin fixed, paraffin embedded and stained with haematoxylin and eosin before tissue histology. Photomicrographs were taken at × 10 or × 20 magnifications.

Statistics. An unpaired two-tailed Student's *t*-test (for two group comparisons) or a two-way analysis of variance (for more than two group comparisons) were performed using Prism (GraphPad) to calculate statistical significance of the difference in mean values and *P* values. A *P* value of <0.05 was considered statistically significant. **P*<0.05; ***P*<0.01. No specific randomization or blinding protocols were used.

Data availability. The data that support the findings of this study are available from the corresponding author on reasonable request. The microarray data have been deposited in the Gene Expression Omnibus under accession GSE97840.

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Author contributions

D.W., Y.L., W.G. and X.F. designed the study; D.W., Y.L., W.G., Q.N., T.X., F.Y. and X.S. performed the experiments and analysed the data; S.C., Y.L., J.L., C.Z. and Z.S. helped with some experiments; H.H., F.L., Z.H., D.Z., Y.Y., G.X., T.C. and X.F. wrote the manuscript; X.F. and T.C. oversaw the project.

Additional information

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