

TSLP links intestinal nutrient sensing with amplification of the ILC2–tuft cell circuit

Received: 19 November 2024

Accepted: 3 October 2025

Published online: 12 November 2025

Chang Liao¹, Elvira Mennillo¹, Michael G. Kattah¹, Ming Ji², Zhi-En Wang², Daniel J. Drucker³, Hong-Erh Liang¹ & Richard M. Locksley^{1,2}✉

Group 2 innate lymphocytes (ILC2s) are prevalent in small intestine but their role during homeostasis is unclear. Here we show that thymic stromal lymphopoietin (TSLP)—a cytokine implicated in ILC2 activation—is expressed constitutively in subepithelial fibroblasts, including telocytes and crypt-associated trophocytes, which are specialized fibroblasts necessary to sustain epithelial identity. Feeding increases TSLP and induces ILC2 type 2 cytokines that are attenuated by deletion of TSLP in fibroblasts or TSLP receptor on ILC2s. Both mouse and human intestinal fibroblasts express receptors for glucagon-like peptide-2 (GLP-2)—an intestinotrophic growth factor released by enteroendocrine cells following food intake. GLP-2 promotes intestinal TSLP in mouse and human intestinal fibroblasts, and TSLP-dependent ILC2 activation and tuft cell hyperplasia in mice, thus linking nutrient detection with ILC2-mediated amplification of the tuft cell chemosensory circuit that promotes epithelial surveillance of ingested cargo.

The small intestinal mucosa represents a key barrier balancing host nutritional needs and protection from potentially noxious chemical and infectious environmental agents. Innate and innate-like lymphocytes, including group 3 innate lymphoid cells, intraepithelial lymphocytes, $\gamma\delta$ T cells, specialized ROR γ ⁺ antigen-presenting cells and regulatory T cells, create a healthy microenvironment that sustains the dynamic interface facilitating nutrient absorption, metabolism, microbiota control, tolerance and epithelial protection during fasting, feeding and circadian cycles^{1–5}. ILC2s are prevalent in small intestinal lamina propria (siLP) although their participation in resting homeostasis is unclear. ILC2s and adaptive Th2 cells become highly engaged during helminth infection and in food allergy but neither of these conditions necessarily explains the basal residence of innate ILC2s in small intestine siLP.

In previous studies, our laboratory and others identified a key role for the alarmin interleukin (IL)-25 as a tuft cell-derived cytokine that activates siLP ILC2s to release IL-13, which biases epithelial progenitors to a secretory cell fate that increases goblet and tuft cells in a forward-amplifying circuit accounting for the ‘weep and sweep’ response to helminths and protists^{6–8}. Deletion of cell intrinsic regulators of IL-25 signaling, such as A20 (encoded by the *Tnfrsf25* gene), resulted in constitutive activation of the circuit, suggesting control

of the basal state⁹. The combinatorial role of alarmins in regulating the activation of tissue resident ILC2s and Th2 cells^{10,11} prompted us to consider roles for additional alarmins in small intestinal physiology and homeostasis. Here we demonstrate that TSLP produced by subepithelial fibroblasts relays signals from epithelial enteroendocrine cells (EECs) to siLP ILC2s to amplify the tuft cell circuit by a reversible process responsive to food intake.

Specialized subepithelial fibroblasts express TSLP in the small intestine

We targeted the *Tslp* locus to identify TSLP-producing cells using the flox-and-reporter (Flare) cassette employed previously to visualize IL-25-producing tuft cells⁶ (Fig. 1a). Under basal conditions in specific pathogen-free C57BL/6 Flare-TSLP mice, the predominant TSLP reporter-positive (TSLP-tandem-dimer red fluorescent protein (tdTomato)⁺) cells in the small intestine were subepithelial PDGFR α ⁺Podoplanin (PDPN)⁺ cells with the morphology of fibroblasts along the villi and enriched at the villus tips throughout small intestine and among crypt-associated cellular networks in the jejunum and ileum (Fig. 1b, Extended Data Fig. 1a and Supplementary Videos 1–7). These cells comprised >80% of TSLP-tdTomato⁺ cells among CD45⁺ cells, with

¹Department of Medicine, University of California, San Francisco, CA, USA. ²Howard Hughes Medical Institute, University of California, San Francisco, CA, USA. ³Lunenfeld-Tanenbaum Research Institute, Mount Sinai Hospital, Toronto, Ontario, Canada. ✉e-mail: richard.locksley@ucsf.edu

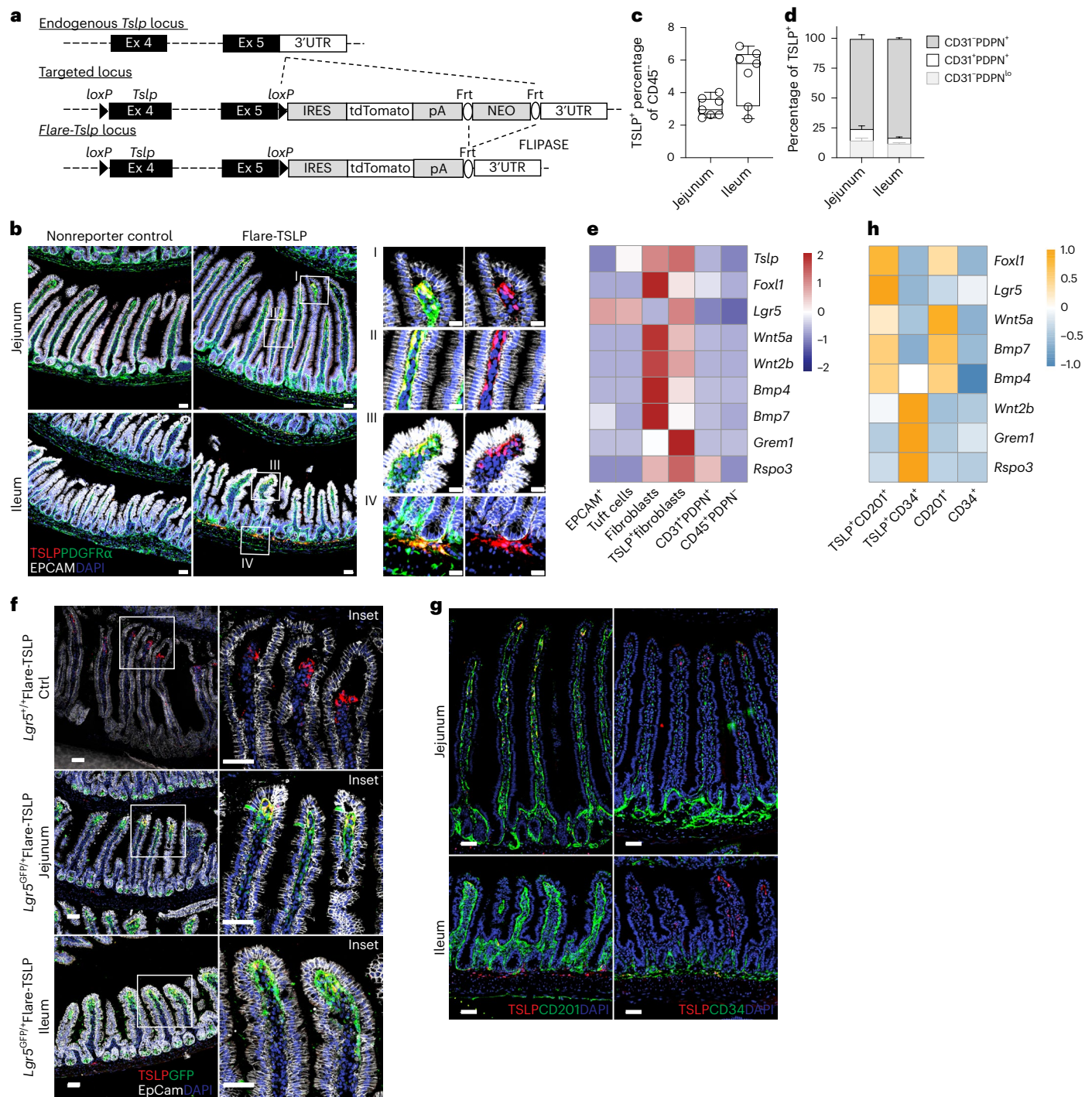


Fig. 1 | Stromal cells are primary sources of TSLP in the small intestine.

a, Gene targeting strategy for the flox-and-reporter of *Tslp* locus. Frt, target site for FLIPASE recombinase; pA, bovine growth hormone poly(A) tail; UTR, untranslated region. **b**, Representative imaging of jejunum (top) and ileum (bottom) of Flare-TSLP or nonreporter control mice. Red, TSLP-tdTomato; green, PDGFRα; white, EPCAM; blue, DAPI; ×20 objective. Scale bars, 50 μm (main images) and 20 μm (boxed areas I–IV). **c, d**, Flow cytometric analysis of stromal populations in jejunum and ileum siLP in Flare-TSLP mice. **c**, Percentage of total TSLP-tdTomato⁺ cells in CD45⁺ population; *N* = 7 biological replicates each column group. **d**, Percentage of endothelial and nonendothelial stromal cells within TSLP-tdTomato⁺ cells. Cells were stained with anti-CD31 and anti-PDPN antibodies; *N* = 3 biological replicates each column group. Error bars indicate samples mean ± s.e.m. **e**, RT-qPCR analysis of sorted CD45⁺ EPCAM⁺ epithelial cells, CD45⁺ EPCAM⁺ Siglec1⁺ tuft cells, CD45⁺ CD31⁺ PDPN⁺ fibroblasts,

CD45⁺ CD31⁺ PDPN⁺ TSLP-tdTomato⁺ fibroblasts, and CD45⁺ CD31⁺ PDPN⁺ lymphatic endothelial and CD45⁺ PDPN⁺ hematopoietic cells from small intestinal epithelial fraction or siLP (whole tissue) of Flare-TSLP mice. Gene expression normalized to 18S. **f**, Representative imaging of jejunum (middle) and ileum (bottom) of Flare-TSLP; *Lgr5*-eGFP dual reporter mice or Flare-TSLP single reporter (Ctrl) mice (top). Note that epithelial tuft cells can express *Lgr5*-eGFP. Red, TSLP-tdTomato; green, *Lgr5*-eGFP; white, EPCAM; blue, DAPI; ×20 objective. Scale bars, 50 μm. **g**, Representative imaging of CD201 and CD34 expression in jejunum (top) and ileum (bottom) of Flare-TSLP mice. Red, TSLP-tdTomato; green, CD201 (left) or CD34 (right); blue, DAPI; ×20 objective. Scale bars, 100 μm. **h**, RT-qPCR analysis of sorted CD45⁺ CD31⁺ TSLP-tdTomato⁺ CD201⁺ CD31⁺, CD45⁺ CD31⁺ TSLP-tdTomato⁺ CD34⁺, CD45⁺ CD31⁺ CD201⁺ and CD45⁺ CD31⁺ CD34⁺ stromal subpopulations from siLP (whole tissue) of Flare-TSLP mice. Gene expression normalized to 18S.

their prevalence increasing from 2–4% in the proximal small intestine to 2–7% in the distal small intestine (Fig. 1c). TSLP-tdTomato⁺CD31⁺ cells were PDPN⁺ LYVE1⁺ lymphatic endothelial cells (LECs) and were less abundant, comprising 10–20% and 1–10% of TSLP-tdTomato⁺CD45⁺ cells in the proximal and distal small intestine, respectively (Fig. 1d and Extended Data Fig. 1b–d). TSLP-tdTomato⁺CD45⁺ cells were rare and found only among the CD31⁺ endothelial cells (Extended Data Fig. 1d). Despite reports of TSLP-positive small intestinal epithelia^{12,13}, including type 2 tuft cells, we did not identify intestinal TSLP-tdTomato⁺ epithelial cells by flow cytometry or microscopy in Flare-TSLP mice (Fig. 1b and Extended Data Fig. 1f). This was not due to an inability to visualize epithelial TSLP, as topical application of the vitamin D analog calcipotriol (MC903), known to induce TSLP in keratinocytes¹⁴, induced reporter signal in Flare-TSLP mice (Extended Data Fig. 1e). In the small intestine, epithelial tuft cells remained TSLP reporter-negative throughout a kinetic analysis of acute *Nippostrongylus brasiliensis* helminth infection (Extended Data Fig. 1f,g). Similarly, proximal large intestine epithelial cells remained TSLP reporter-negative after *Trichuris muris* helminth infection, which is known to require TSLP¹² (Extended Data Fig. 1h). After both infections, however, reporter expression was present in PDGFR α ⁺ subepithelial fibroblasts (Extended Data Fig. 1g,h).

The location and morphology of constitutive TSLP-tdTomato⁺ subepithelial fibroblasts in Flare-TSLP mice resembled previous reports of telocytes and crypt-associated trophocytes. These structural cells generate the counter-regulating WNT and BMP gradients that maintain the crypt stem cell niche and regulate epithelial differentiation during vertical zonation of the villi^{15,16}. To confirm the identity of these cells, we employed fluorescence-activated cell sorting (FACS) to enrich TSLP-tdTomato⁺ cells and used quantitative PCR (qPCR) with reverse transcription (RT) (RT-qPCR) to assess the expression of signature transcripts for telocytes, including *Foxl1*, *Wnt5a* and *Bmp7*, and trophocytes, including *Wnt2b*, *Grem1* and *Rspo3* (refs. 15,16; Fig. 1e). Expression of these genes was relatively low in epithelial cell adhesion molecule (EPCAM)⁺ total epithelial cells, tuft cells, CD45⁺CD31⁺PDPN⁺ LECs or CD45⁺PDPN⁺ hematopoietic cells, with the exception of *Rspo3*, which was also highly expressed by LECs (Fig. 1e). The expression of *Lgr5* in the TSLP-tdTomato⁺ population (Fig. 1e) suggested that villus tip telocytes (VTTs), specialized telocytes that act as signaling hubs to direct apical epithelial cell fate¹⁷, also express TSLP. Using Flare-TSLP;*Lgr5*-eGFP dual reporter mice, we confirmed co-expression of *Tslp* and *Lgr5* among VTTs (Fig. 1f). Previous studies identified CD201 and CD34 as markers for intestinal telocytes and trophocytes, respectively^{16,18}. In line with this, CD201 labeled villus subepithelial fibroblasts and *Lgr5*-GFP⁺ VTTs but not trophocytes, while CD34 labeled crypt-associated TSLP-tdTomato⁺ trophocytes (Fig. 1g and Extended Data Fig. 2a). We sorted CD45⁺CD31⁺CD201⁺TSLP-tdTomato⁺ and CD45⁺CD31⁺CD34⁺TSLP-tdTomato⁺ cells (Extended Data Fig. 2b) and confirmed their expression of telocyte- and trophocyte-associated transcripts, respectively (Fig. 1h and Extended Data Fig. 2c). Thus, distinct subepithelial fibroblasts, including villus telocytes and trophocytes, as well as LECs, constitute the primary cell types that express TSLP in the mouse small intestine under resting conditions.

TSLP increases with feeding and activates ILC2 cytokine production

Feeding activates type 2 cytokine expression by small intestine ILC2s independently of the light–dark cycle¹⁹. After 16-h fasting, Flare-TSLP mice were orally gavaged with a 500- μ l slurry of powdered diet containing ~2 kcal mixed nutrients (about 1% total daily caloric intake with components proportional to standard chow diet) or water control (Fig. 2a). After 2 h, the prevalence of TSLP-tdTomato⁺CD45⁺ nonepithelial cells increased in proximal and distal siLP (Fig. 2b and Extended Data Fig. 3a). We confirmed TSLP protein upregulation by enzyme-linked immunosorbent assay (ELISA) in supernatants from small intestine tissue

explants and homogenates, observing that TSLP amounts increased and peaked around 2 hr after gavage (Fig. 2c and Extended Data Fig. 3b,c). Notably, TSLP upregulation at this timepoint was unchanged after nutrient gavage of fasted *Pou2f3*^{−/−} mice, which lack tuft cells, indicating that tuft cells were not required for this increase (Fig. 2c and Extended Data Fig. 3d,e). To investigate the role for TSLP in ILC2 activation after feeding, we utilized cytokine reporter mice²⁰ with marker alleles for IL-5 (Red5) and IL-13 (Smart13) crossed to TSLPR-deficient (*Tslpr*^{−/−}) mice. We analyzed ILC2s from proximal siLP after 16-h fasting and 4-h refeeding with chow diet and water ad libitum or water alone (Fig. 2d). Consistent with previous findings¹⁹, ILC2s were activated 4 h after feeding, as evidenced by an increase in proportion of IL-13⁺ (Smart13)⁺ ILC2s that was attenuated in TSLPR-deficient mice (Fig. 2e,f). Furthermore, we generated ILC2 deletion of the TSLP receptor by crossing *Tslpr*^{fl/fl} mice with *IL5*^{Cre+} mice and confirmed that the loss of TSLPR expression on ILC2s impaired their activation after feeding (Fig. 2g).

We next sought to show the contributions of stromal-derived TSLP to activation of intestinal ILC2s by food intake. To achieve this, we generated *Tslp*^{fl/fl}*Pdgfra*^{CreERT2}Smart13 and *Tslp*^{fl/fl}*Lyve1*^{Cre}Smart13 reporter mice, which delete TSLP from fibroblasts and LECs, respectively. Following administration of tamoxifen to *Tslp*^{fl/fl}*Pdgfra*^{CreERT2}Smart13 mice, the loss of TSLP from fibroblasts resulted in attenuation of ILC2 activation after feeding, as indicated by diminished expression of the IL-13 reporter (Fig. 2h,i). In contrast, ILC2 activation in *Tslp*^{fl/fl}*Lyve1*^{Cre}Smart13 mice remained intact (Fig. 2j). Thus, fibroblasts, rather than LECs, are the primary source of TSLP regulating ILC2 activation after feeding.

GLP-2 mediates upregulation of TSLP in subepithelial fibroblasts

To better understand the cells underlying feeding-induced TSLP-ILC2 signaling, we performed single-cell RNA sequencing (scRNA-seq) of TSLP-tdTomato⁺ cells isolated from the proximal and distal siLP of Flare-TSLP mice using at least three biological samples from each tissue (Extended Data Fig. 4a). Unsupervised cell clustering analysis revealed the main stromal cell types were fibroblasts and LECs, and consistent with the flow cytometric analyses (Fig. 3a and Extended Data Figs. 4b and 5a). Principal subsets of fibroblasts included subepithelial myofibroblasts (SEMFs) (which include *Foxl1*⁺ telocytes), *Lgr5*⁺ VTTs, *Mik67*⁺ proliferative telocytes, *Grem1*⁺ trophocytes and recently described *Pil6*⁺ ‘universal’ fibroblasts with transcriptomic signatures of mesothelial-like cells²¹ (Fig. 3a and Extended Data Fig. 4b,c). In accordance with their anatomic locations, SEMFs and trophocytes were highlighted by expression of distinct WNTs and BMPs (Extended Data Fig. 4d). SEMFs were enriched for noncanonical WNTs (such as *Wnt4* and *Wnt5a*) and *Bmps*, and typically expressed by telocytes. Notably, *Bmp7* is specifically enriched in VTTs. In contrast, trophocytes were enriched for canonical WNTs, such as *Wnt2b*, and BMP inhibitors, including *Grem1*, *Rspo3* and *Dkk2*. The three trophocyte subsets exhibited similar transcriptomic signatures, as did SEMFs, VTTs and proliferative telocytes (Extended Data Fig. 4e). These cell types expressed comparable gene signatures across the proximal and distal regions of small bowel (Extended Data Figs. 4e and 5c).

Nutrients in food are absorbed primarily by the prevalent enterocytes and sensed by less frequent EECs, which respond to luminal content by releasing peptide hormones that regulate nutrient uptake, metabolism, intestinal motility and satiety. EECs, named by dominant hormones they release, develop along vertically zoned villus trajectories through nodes of differentiation accompanied by activation of overlapping components of their peptide hormone repertoires^{22,23}. We next investigated the expression of gut peptide hormone receptors expressed by subepithelial fibroblasts that would enable crosstalk between nutrient-sensing EECs and these structural cells. Our data, consistent with that of others, revealed that intestinal stromal cells express high amounts of mRNA for the receptor for glucagon-like peptide-2 (GLP-2) with less consistent evidence for other gut peptide

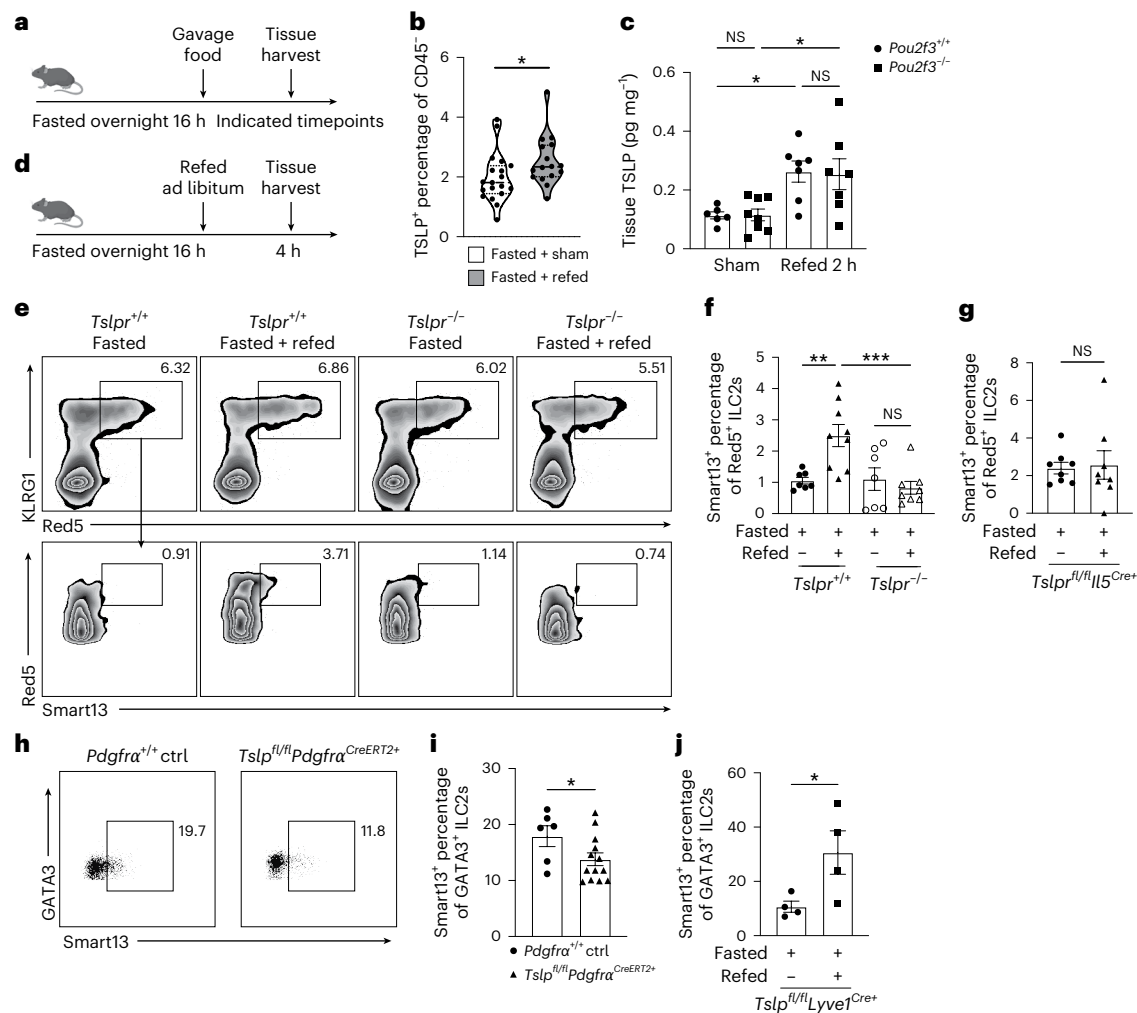


Fig. 2 | Feeding increases intestinal TSLP and drives ILC2 activation.

a, Schematic of the protocol for measuring intestinal TSLP after feeding. Mice were fasted for 16 h overnight before oral gavage with 500 μ l food slurry (refed) or water as volumetric control (sham). Tissues were harvested as indicated. **b**, Percentage of TSLP-tdTomato⁺ cells among CD45⁺ cells in proximal jejunal siLP by flow cytometric analysis after oral gavage at 2 h after overnight fasting. Biological replicates $N = 19$ for sham control group and $N = 16$ for refeeding group. Statistical analysis was performed using unpaired t -test, $*P < 0.05$. Error bars indicate samples mean \pm s.e.m. **c**, ELISA of TSLP protein from supernatant of proximal jejunal tissue explants after oral gavage at 2 h in *Pou2f3*^{-/-} or WT mice after overnight fasting. Biological replicates with total $N = 28$, pooled from 2 independent experiments. Statistical analysis was performed using ANOVA with correction for multiple comparisons, $*P < 0.05$. Error bars indicate samples mean \pm s.e.m. NS, nonsignificant. **d**, Schematic of the protocol for measuring siLP ILC2 activation after feeding. Mice were fasted for 16 h overnight and given access to standard chow and water ad libitum (refed) or maintained on water control (fasted). Tissues were harvested 4 h later. **e, f**, Percentage of IL-13 (Smart13)⁺ ILC2s among ILC2s in proximal jejunal LP in *Tslpr*^{+/+}Red5Smart13 or *Tslpr*^{-/-}Red5Smart13 mice after 16-h fasting followed by 4-h refeeding ad libitum. **e**, Representative flowplots. **f**, Quantification. ILC2s were gated as Lin⁻CD45⁺IL-5 (Red5)⁺ cells. Biological replicates with total $N = 31$, pooled from at least 2

independent experiments. Statistical analysis was performed using ANOVA with correction for multiple comparisons, $**P < 0.01$, $***P < 0.001$. Error bars indicate samples mean \pm s.e.m. **g**, Percentage of IL-13 (Smart13)⁺ ILC2s among ILC2s in proximal jejunal LP in *Tslpr*^{fl/fl}IL5^{Cre+} (Red5)Smart13 mice after 16-h fasting followed by 4-h water or refeeding ad libitum. ILC2s were gated as Lin⁻CD45⁺IL-5 (Red5)⁺ cells. Biological replicates $N = 8$ each column group, pooled from 2 independent experiments. Statistical analysis was performed using unpaired t -test. Error bars indicate samples mean \pm s.e.m. **h, i**, Percentage of IL-13 (Smart13)⁺ ILC2s among total ILC2s in proximal jejunal LP 4 h after refeeding ad libitum in overnight fasted *Tslpr*^{fl/fl}*Pdgfra*^{CreERT2+}Smart13 or littermate control *Pdgfra*^{+/+}Smart13 mice (ctrl) post-tamoxifen. **h**, Representative flowplots. **i**, Quantitation. ILC2s were gated on Lin⁻CD45⁺GATA3⁺ cells. Biological replicates $N = 6$ for control group and $N = 13$ for experiment group, pooled from at least 2 independent experiments. Statistical analysis was performed using unpaired t -test, $*P < 0.05$. Error bars indicate samples mean \pm s.e.m. **j**, Percentage of IL-13 (Smart13)⁺ ILC2s among ILC2s in proximal jejunal LP 4 h after water or refeeding ad libitum in overnight fasted *Tslpr*^{fl/fl}*Lyve*^{Cre+}Smart13 mice. ILC2s were gated on Lin⁻CD45⁺GATA3⁺ cells. Biological replicates $N = 4$ in each column group. Statistical analysis was performed using unpaired t -test, $*P < 0.05$. Error bars indicate samples mean \pm s.e.m. Illustrations in **a** created with BioRender.com.

hormone receptors (Fig. 3b,c), in both mouse and human^{15,16,24,25} (Extended Data Fig. 6a,e–h). Spatial transcriptional analysis from a public dataset of mouse small intestine revealed that *Glp2r* expression associated predominantly with *Pdgfra*^{hi} populations that also co-expressed *Tslp* (Extended Data Fig. 6b–d). Of note, *Foxl1*⁺ telocytes, including VTTs, were enriched strongly for *Glp2r* expression (Fig. 3b and Extended Data Fig. 4f).

Although prominent in pancreatic glucagon-producing alpha cells, proglucagon is also expressed highly in subsets of small intestinal EECs, particularly L cells, which express tissue-specific convertases that post-translationally process proglucagon to the peptides glucagon-like peptide-1 (GLP-1), GLP-2 and oxyntomodulin rather than glucagon^{26–28}. In response to food constituents, proglucagon-derived peptides are secreted into the basolateral space, where the activities of GLP-1 and

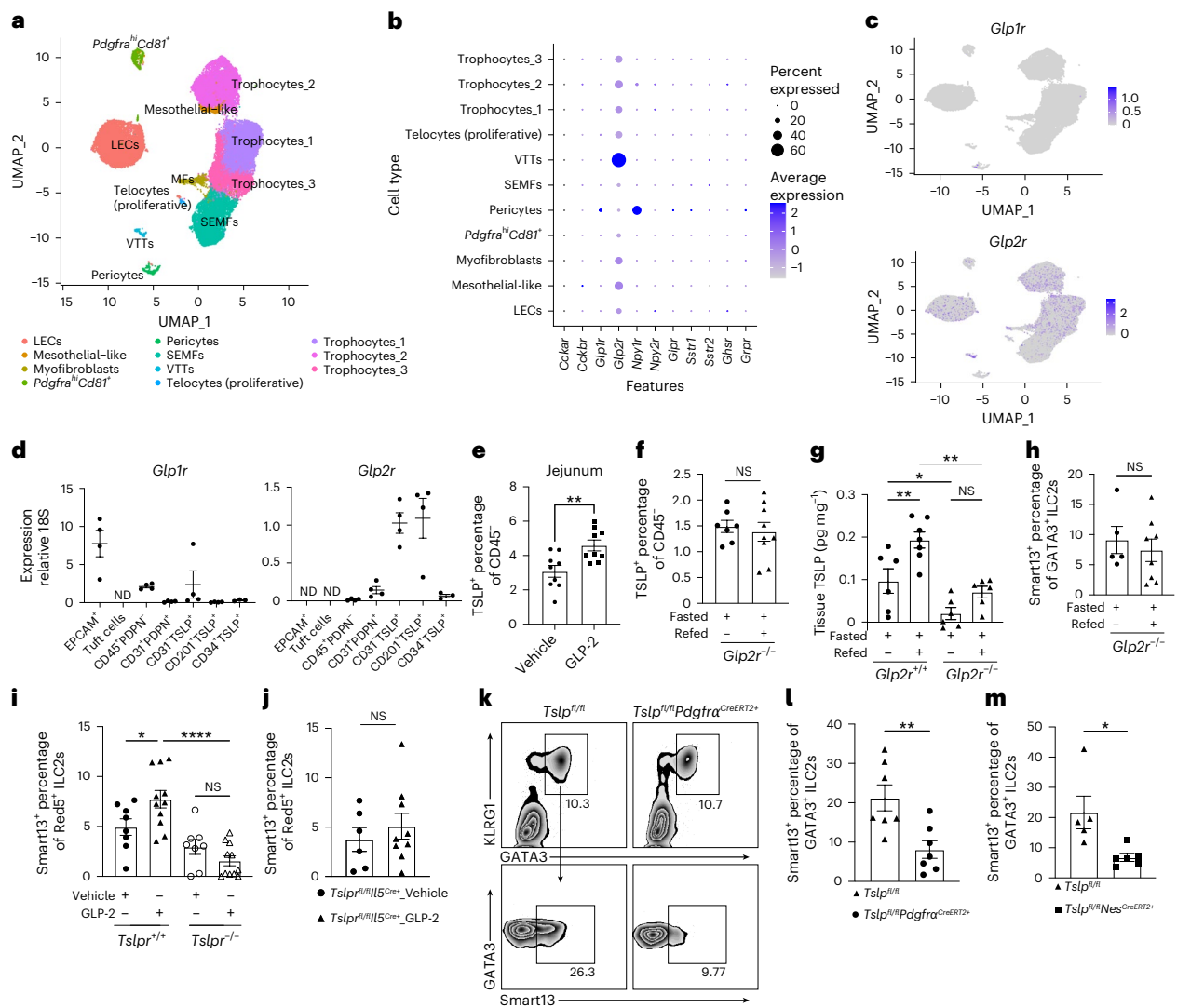


Fig. 3 | GLP-2 drives ILC2 activation through TSLP signaling. a–c, scRNA-seq analysis of sorted TSLP-tdTomato⁺ cells from the siLP. **a**, UMAP representing cell clustering analysis. **b**, Expression of gut hormone receptors by TSLP-tdTomato⁺ stromal cells. **c**, Feature plots representing *Glp1r* (top) and *Glp2r* (bottom) expression. **d**, RT-qPCR analysis of *Glp1r* (left) and *Glp2r* (right) expression on sorted gut. CD45⁺EPCAM⁺ (epithelial cells), tuft cells, CD45⁺PDPN⁺ (hematopoietic cells), CD45⁺CD31⁺PDPN⁺ (LECs), CD45⁺CD31⁺TSLP-tdTomato⁺ total stromal cells, CD45⁺CD31⁺TSLP-tdTomato⁺CD201⁺ (telocytes) and CD45⁺CD31⁺TSLP-tdTomato⁺CD34⁺ (trophocytes). Biological replicates $N = 3$ or 4, pooled from 2 independent experiments. Error bars indicate samples mean \pm s.e.m. **e**, Percentage of TSLP-tdTomato⁺ cells among CD45⁺ cells in proximal jejunal LP after three consecutive GLP-2[Gly2] s.c. injections as assessed by flow cytometric analysis. Biological replicates $N = 9$ for each column group, pooled from at least 2 independent experiments. Statistical analysis was performed using unpaired t -test, $^{**}P < 0.01$. Error bars indicate samples mean \pm s.e.m. **f**, Flow cytometric analysis showing percentage of total TSLP-tdTomato⁺ cells among CD45⁺ cells in proximal jejunal LP from fasted *Glp2r*^{-/-} Flare-TSLP mice after oral food gavage at 2 h. Biological replicates $N = 7$ for control group and $N = 9$ for refeeding group, pooled from at least 2 independent experiments. Statistical analysis was performed using unpaired t -test. Error bars indicate samples mean \pm s.e.m. **g**, ELISA of TSLP protein from supernatant of proximal jejunal tissue explants after oral gavage at 2 h in *Glp2r*^{-/-} or WT mice after overnight fasting. Biological replicates with total $N = 25$, pooled from at least 2 independent experiments. Statistical analysis was performed using ANOVA with correction for multiple comparisons, $^{*}P < 0.05$, $^{****}P < 0.0001$. Error bars indicate samples mean \pm s.e.m. **h**, Quantification of percentage of IL-13 (Smart13⁺) ILC2s among proximal jejunal LP ILC2s from fasted *Glp2r*^{-/-} Smart13 mice after 16-h fasting followed by 4-h water or refeeding ad libitum. ILC2s were

gated on Lin⁺CD45⁺GATA3⁺ cells. Biological replicates $N = 5$ for control group and $N = 8$ for refeeding group, pooled from at least 2 independent experiments. Statistical analysis was performed using unpaired t -test. Error bars indicate samples mean \pm s.e.m. **i**, Quantification of percentage of IL-13 (Smart13⁺) ILC2s among proximal jejunal LP ILC2s in *Tslpr*^{+/+} Red5Smart13 or *Tslpr*^{-/-} Red5Smart13 mice after three daily injections of GLP-2[Gly2]. ILC2s were gated on Lin⁺CD45⁺IL-5(Red5⁺) cells. Biological replicates with total $N = 38$, pooled from at least 2 independent experiments. Statistical analysis was performed using ANOVA with correction for multiple comparisons, $^{*}P < 0.05$, $^{****}P < 0.0001$. Error bars indicate samples mean \pm s.e.m. **j**, Quantification of percentage of IL-13 (Smart13⁺) ILC2s among proximal jejunal LP ILC2s in *Tslpr*^{+/+} IL5^{Cre+} Smart13 after three daily injections of GLP-2[Gly2] or vehicle control. ILC2s were gated on Lin⁺CD45⁺IL-5(Red5⁺) cells. Biological replicates $N = 6$ for vehicle control group and $N = 9$ for GLP-2[Gly2] injection group, pooled from 2 independent experiments. Statistical analysis was performed using unpaired t -test. Error bars indicate samples mean \pm s.e.m. **k, l**, Flow cytometric analysis of percentage of IL-13 (Smart13⁺) ILC2s among proximal jejunal LP ILC2s in *Tslpr*^{+/+} *Pdgfra*^{CreERT2+} Smart13 or littermate *Tslpr*^{+/+} Smart13 controls post-tamoxifen followed by three daily injections of GLP-2[Gly2]. **k**, Representative flowplots. **l**, Quantitation. ILC2s were gated as Lin⁺CD45⁺GATA3⁺ cells. Biological replicates $N = 7$ for each column group, pooled from at least 2 independent experiments. Statistical analysis was performed using unpaired t -test, $^{**}P < 0.01$. Error bars indicate samples mean \pm s.e.m. **m**, Flow cytometric analysis of percentage of IL-13 (Smart13⁺) ILC2s among proximal jejunal LP ILC2s in *Tslpr*^{+/+} *Nes*^{CreERT2+} Smart13 or littermate *Tslpr*^{+/+} Smart13 controls post-tamoxifen followed by three daily injections of GLP-2[Gly2]. Biological replicates $N = 6$ for control group and $N = 5$ for experiment group, pooled from 2 independent experiments. Statistical analysis was performed using unpaired t -test, $^{*}P < 0.05$. Error bars indicate samples mean \pm s.e.m.

GLP-2 are kept localized by high concentrations of the inactivating dipeptidyl peptidase-4 enzyme. GLP-2 is intestinotrophic, promoting microvillus lengthening, increasing vascular flow and enhancing the epithelial barrier, in part by stimulating release of insulin-like growth factor-1 and epidermal growth factor from subepithelial fibroblasts²⁶. Using qPCR analysis, we detected *Glp2r* expression by stromal cells but not epithelial cells (Fig. 3d). Further, administration of stabilized GLP-2R agonist, GLP-2[Gly2], significantly increased the prevalence of TSLP-tdTomato⁺ cells in siLP compared to mice treated with vehicle control (Fig. 3e and Extended Data Fig. 7a,b). The proportions of fibroblasts or LECs were not altered by GLP-2[Gly2] administration (Extended Data Fig. 7c), but there appeared to be more CD201⁺ telocytes within the CD45⁺TSLP-tdTomato⁺CD31⁺ stromal populations in the siLP (Extended Data Fig. 7d,e). In vitro incubation of sorted CD45⁺TSLP-tdTomato⁺CD31⁺ cells with GLP-2[Gly2] resulted in increased antibody labeling of TSLP protein compared to vehicle control-treated cells (Extended Data Fig. 7f).

In considering how GLP-2 might induce TSLP expression in the gut subepithelial fibroblasts, we note previous studies implicating GLP-2 stimulation of enteric neurons by signals through PI3K with mTOR or ERK to induce VIP expression²⁹. GLP-2 can also increase intracellular cyclic AMP concentrations, including in fibroblasts^{30,31}, and cAMP has been implicated in TSLP production and release^{32,33}. To investigate the potential mechanisms by which GLP-2 regulates TSLP upregulation, we measured TSLP protein concentrations in three experimental settings, including primary intestinal fibroblasts derived from mouse and human tissues as well as ex vivo mouse intestinal explants. Fibroblast monolayers or tissue explants were treated with GLP-2 alone or in combination with a cAMP agonist forskolin or with inhibitors targeting CREB (666-15), PI3K (LY294002), mTOR (rapamycin), or MEK/ERK (PD98059) (Extended Data Fig. 7g–j). Treatment with GLP-2 agonist alone significantly increased TSLP amounts, whereas cotreatment with PI3K, mTOR and MEK/ERK inhibitors blocked the increased TSLP in all three models, suggesting the engagement of PI3K–mTOR and PI3K–ERK signaling pathways, and consistent with previous findings. In mouse and human fibroblasts, forskolin increased TSLP and antagonism of CREB using 666-15 abrogated GLP-2-induced TSLP upregulation (Extended Data Fig. 7g,h,j); mouse explants were less affected by interruption of these pathways (Extended Data Fig. 7i). Together, these data suggest that GLP-2 regulates TSLP upregulation through PI3K–mTOR and PI3K–ERK signaling with the possible involvement of the cAMP–CREB axis in TSLP induction.

GLP-2 mediates TSLP-dependent activation of intestinal ILC2s in vivo

In support of the role for GLP-2 in the induction of TSLP by feeding, the increased prevalence of TSLP-tdTomato⁺ cells in proximal siLP and the amounts of TSLP protein recovered from intestinal tissue explants and homogenates were abrogated in fasted and refed mice lacking GLP-2R (Fig. 3f,g and Extended Data Fig. 7k). Activation of ILC2s after feeding was also attenuated in *Glp2r*^{−/−} mice as assessed by IL-13 reporter (Smart13) expression (Fig. 3h). These results suggest that both TSLP production and ILC2 activation are downstream effects of GLP-2 induced by feeding. Administration of GLP-2[Gly2] led to an increase in activated IL-13-reporter⁺ ILC2s that was not associated with proliferation and that was attenuated in TSLPR-deficient mice (Fig. 3i and Extended Data Fig. 7l,m), as well as in *Tslpr*^{fl/fl}IL5^{Cre+} Smart13 mice (Fig. 3j), which lack TSLPR on ILC2s, and tamoxifen-treated *Tslpr*^{fl/fl} *Pdgfra*^{CreERT2+} Smart13 mice (Fig. 3k,l), which lack TSLP in fibroblasts.

TSLP is also important for type 2 immune responses after *T. muris* infection¹². In line with this, the activation of *Tslpr*^{−/−} ILC2s was diminished significantly following *T. muris* infection (Extended Data Fig. 8a). A similar reduction in ILC2 activation was observed in *Tslpr*^{fl/fl}IL5^{Cre+} Smart13 mice (Extended Data Fig. 8b), and *Tslpr*^{fl/fl} *Pdgfra*^{CreERT2+} Smart13 after tamoxifen treatment (Extended Data Fig. 8c).

Given that *Glp2r* expression was highly expressed in telocytes among stromal cell populations in the small intestine (Fig. 3b and Extended Data Fig. 5b), we hypothesize that these cells may represent the primary GLP-2-responsive cells mediating the downstream TSLP–ILC2 signaling axis. qPCR analysis revealed that *Nes*, the gene encoding for the intermediate filament protein Nestin, was expressed by total stromal cells, with CD201⁺ subepithelial villus fibroblasts including telocytes showing the highest amounts of *Nes* expression as compared to other stromal subsets, including CD34⁺ trophocytes (Extended Data Fig. 8d). We crossed *Tslpr*^{fl/fl} Smart13 reporter mice with *Nes*^{CreERT2} mice to achieve deletion of *Tslp* in subepithelial villus fibroblasts. After tamoxifen treatment, *Tslpr*^{fl/fl} *Nes*^{CreERT2+} Smart13 mice exhibited diminished ILC2 activation following GLP-2[Gly2] administration and this was phenocopied using *Tslpr*^{fl/fl} *Pdgfra*^{CreERT2+} Smart13 mice (Fig. 3m), indicating that subepithelial villus fibroblasts represent the main source of TSLP in the small intestine that mediates ILC2 activation downstream of GLP-2. Further, after *T. muris* infection, *Tslpr*^{fl/fl} *Nes*^{CreERT2+} Smart13 mice also exhibited impaired ILC2 activation as indicated by the reduced percentage of Smart13⁺ ILC2s and lower expression of Smart13 and ICOS (Extended Data Fig. 8e,f), and consistent with enriched *Glp2r* expression by TSLP-tdTomato⁺ subepithelial fibroblasts, including telocytes, in the mouse cecum and proximal colon (Extended Data Fig. 5b) where the parasites reside. These data indicate that fibroblast-derived TSLP, particularly from the telocytes, is required for optimal activation of TSLPR⁺ ILC2s after food intake and *T. muris* infection. Of note, male *Glp2r*^{−/−} mice had diminished ILC2 activation compared to wild-type (WT) mice following *T. muris* infection, whereas female *Glp2r*^{−/−} mice were more variable (Extended Data Fig. 8g,h), and suggesting that GLP-2 may play a role in TSLP-dependent ILC2 activation during *T. muris* infection.

Taken together, these data support a biological pathway linking the activation of preproglucagon⁺ EECs with GLP-2-mediated stimulation of subepithelial fibroblasts leading to increased production of TSLP, which activates resident ILC2s to produce IL-13 in the intestine.

GLP-2 mediates ILC2-dependent amplification of the small intestinal tuft cell circuit

Activation of intestinal ILC2s to release IL-13 leads to biased production of tuft cells from the transit-amplifying zone, which depends on IL-4Rα expression on intestinal epithelial cells^{6,7}. Consistent with these findings, administration of GLP-2[Gly2] resulted in increased numbers of small intestinal tuft cells independent of proliferation as indicated by 5-ethynyl-2'-deoxyuridine (EdU) negative staining, which was significantly attenuated in *Tslpr*^{−/−}, *Tslpr*^{fl/fl}IL5^{Cre+}, *Il13ra1*^{−/−} and *Il4ra*^{−/−} mice (Fig. 4a,b and Extended Data Fig. 9a,b). Further, the increase in tuft cells was diminished in *Il4ra*^{fl/fl} *Vil1*^{Cre+} mice, supporting a role for epithelial IL-4Rα in activating the circuit (Fig. 4b and Extended Data Fig. 9a,b). Furthermore, GLP-2-induced tuft cell hyperplasia was attenuated in tamoxifen-treated *Tslpr*^{fl/fl} *Pdgfra*^{CreERT2+} and *Tslpr*^{fl/fl} *Nes*^{CreERT2+} mice that lack fibroblast TSLP as compared to control mice (Fig. 4c and Extended Data Fig. 9c,d). To further confirm EECs can activate the ILC2–tuft cell circuit through an activated G protein-coupled receptor (GPCR) pathway, we utilized triple-mutant *Vil1*^{Flp} *Cck*^{Cre} *R26*^{Dual-hm3Dq} mice with epithelial lineage-restricted expression of activating designer receptors exclusively activated by designer drugs (DREADDs) on CCK-lineage-restricted intestinal epithelia, which includes preproglucagon⁺ EECs that secrete GLP-2 (ref. 23; Fig. 4d and Extended Data Fig. 9e). Following administration of the synthetic ligand clozapine N-oxide (CNO), ILC2s and tuft cells were increased in the small intestine, confirming amplification of the epithelial tuft cell circuit downstream of EEC activation (Fig. 4e–g and Extended Data Fig. 9f,g).

Discussion

Intestinal telocytes and trophocytes are subepithelial fibroblasts extending from the crypts to the villus tips whose arborizing dendrites

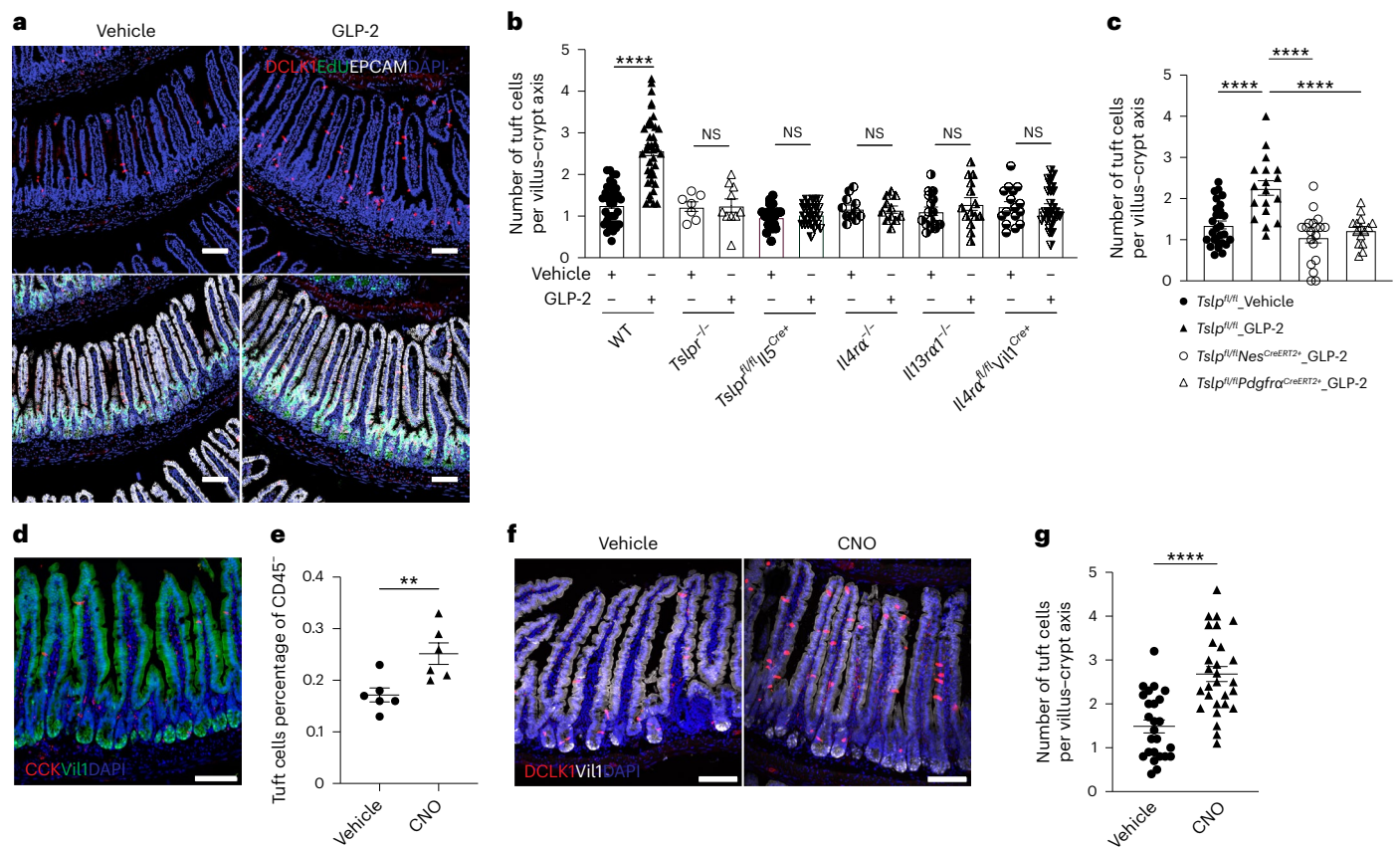


Fig. 4 | GLP-2 drives ILC2-dependent tuft cell expansion. **a**, Representative imaging of tuft cells in jejunum after three daily injections of GLP-2[Gly2]. EdU was injected 24 h before tissue harvest. Green, EdU; red, DCLK1; white, EPCAM; blue, DAPI; $\times 20$ objective. Scale bars, 100 μ m. **b**, Quantification of jejunal tuft cells after three daily GLP-2[Gly2] injections or vehicle control in WT, *Tslpr*^{-/-}, *Tslpr*^{fl/fl}*Il5*^{Cre+}, *Il4ra*^{-/-}, *Il13ra1*^{-/-} and *Il4ra*^{fl/fl}*Vil1*^{Cre+} mice. For tuft cell quantification, images were acquired using $\times 20$ objective, and total DCLK1⁺EPCAM⁺ cells in each field were counted and normalized by total number of villus–crypt axes in the field. Each dot represents an imaging field, total $N = 232$; data pooled from 3 to 10 mice. A plot with averages of each mouse is included in Extended Data Fig. 9b. Statistical analysis was performed using ANOVA with correction for multiple comparisons, **** $P < 0.0001$. Error bars indicate samples mean \pm s.e.m. **c**, Quantification of jejunal tuft cells after three daily GLP-2[Gly2] or vehicle control injections in *Tslpr*^{fl/fl}, *Tslpr*^{fl/fl}*Nes*^{CreERT2+} and *Tslpr*^{fl/fl}*Pdgfra*^{CreERT2+} mice. Each dot represents an imaging field, total $N = 80$; data pooled from 3 to 6 mice. A plot with averages of each mouse is included in Extended Data Fig. 9d. Statistical analysis was performed using ANOVA with correction for multiple comparisons,

**** $P < 0.0001$. Error bars indicate samples mean \pm s.e.m. **d**, Representative imaging showing CCK⁺ EECs in *Vil1*^{fl/fl}*Cck*^{Cre}*R26*^{Dual-HM3Dq} mice. Red, Cck-mCherry; green, Vil1-GFP; blue, DAPI; $\times 20$ objective. Scale bar, 100 μ m. **e–g**, Quantification of jejunal tuft cells after administration of clozapine N-oxide CNO or vehicle control in *Vil1*^{fl/fl}*Cck*^{Cre}*R26*^{Dual-HM3Dq} mice. **e**, Percentage of tuft cells among the CD45⁺ epithelial fraction by flow cytometric analysis. Two independent experiment repeats were combined to generate the plot. Biological replicates $N = 6$ for each column group were compared using two-tailed unpaired t -test, ** $P < 0.005$. Error bars indicate samples mean \pm s.e.m. **f**, Representative imaging of tuft cells. Red, DCLK1; white, Vil1-GFP; blue, DAPI; $\times 20$ objective. Scale bars, 100 μ m. **g**, Quantification of tuft cells. Each dot represents an imaging field, $N = 25$ for control group and $N = 28$ for CNO treatment group; data pooled from 6 mice each group from 2 independent experiments. A plot with averages of each mouse is included in Extended Data Fig. 9g. Statistical analysis was performed using two-tailed unpaired t -test, **** $P < 0.0001$. Error bars indicate samples mean \pm s.e.m.

establish the critical WNT and BMP counter-gradients that maintain stem cells and control epithelial differentiation during villus transit^{15,16,34,35}. Our findings demonstrate that these critical structural cells are the principal intestinal cell sources of TSLP—an alarmin linked with activation of type 2 immune cells, including ILC2s—and are consistent with roles for long-lived structural cells as hubs for tissue immunity³⁶. Notably, TSLP expression was prominent in VTTs, specialized *Lgr5*⁺ subepithelial fibroblasts that control epithelial differentiation at villus apices, and positioning them to relay information regarding intestinal content¹⁷. Further work is needed to assess whether related structural fibroblast subsets in other tissues mediate local TSLP-mediated communication with innate immune cells.

Although activation of ILC2s by food intake was noted¹⁹, how information regarding luminal content is relayed through epithelia to resident ILC2s remained unclear. EECs sense nutrients and secrete peptide hormones and neurotransmitters that collectively orchestrate

intestinal motility, absorption and integration of neuronal circuitry controlling feeding, satiety and positive consumptive or negative aversive reactions³⁷. Sophisticated fate-mapping and lineage-marking studies revealed EECs have a longer half-life than enterocytes during villus ascent and are zoned vertically such that peptide hormones are expressed sequentially in overlapping patterns among EEC subsets to optimize small intestinal function^{22,23,38}. In our investigation, we verified expression of receptors on telocytes for GLP-2—a proglucagon-derived peptide previously shown to promote release of epithelial growth factors from intestinal subepithelial fibroblasts³⁹. Although GLP-2 deletion does not impact small intestine differentiation, it is required for recovery from intestinal atrophy after fasting or injury^{40–42}, underpinning the success of stabilized GLP-2 receptor agonists used therapeutically in humans to increase small intestine growth or prevent atrophy following parenteral feeding²⁷. Although effects of GLP-2 receptor agonists are reversible after cessation of therapy⁴³, the striking intestinotrophic

effects of model helminth and protist infections prompted us to query whether this physiologic pathway shared mechanisms with the intestinal response to these organisms, which includes amplification of the ILC2–tuft cell circuit. Whether GLP-1, which is generated in equimolar amounts with GLP-2 after nutrient ingestion, is involved in metabolic adaptations induced by helminth infection and innate type 2 immune pathways will require further study^{9,27,28,44}.

Our findings in mouse and human systems show that GLP-2 can stimulate telocytes to produce TSLP, thus linking nutrient ingestion and ILC2-mediated amplification of epithelial tuft cells. Tuft cells are specialized chemosensory cells that express a variety of GPCRs that share signal transduction recognition with type 2 taste receptors and provide feedback amplification of the ILC2–tuft cell circuit by release of a second alarmin, IL-25 (ref. 45). As suggested recently, such a system may be part of a regulated food quality control circuit by increasing detection of constituents in ingested cargo that attach positive or negative associations on content that has passed more proximal surveillance mechanisms⁴⁶. Although nutrients stimulate EECs and indirectly activate reversible tuft cell expansion through GLP-2 and TSLP, intestinal protists and helminths can stimulate tuft cell GPCRs directly and sustain the amplified circuit^{9,47,48}. By bypassing a physiologic circuit linked with feeding, endemically adapted parasites establish a reproductive niche protected from superinfection by other pathogens while sustaining the host metabolic state^{9,49,50}—a condition supported by intestinal lengthening, tuft cell expansion and increased attentiveness to ingested constituents (Extended Data Fig. 9h). Understanding how alarmins like TSLP and IL-25 act combinatorially to regulate innate type 2 immunity in support of local physiologic pathways, as co-opted by intestinal pathobionts to sustain their niche, may enable new strategies for increasing tissue resilience while avoiding the dysregulated state of allergic disease.

Online content

Any methods, additional references, Nature Portfolio 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/s41590-025-02328-y>.

References

- Godinho-Silva, C. et al. Light-entrained and brain-tuned circadian circuits regulate ILC3s and gut homeostasis. *Nature* **574**, 254–258 (2019).
- Sullivan, Z. A. et al. $\gamma\delta$ T cells regulate the intestinal response to nutrient sensing. *Science* **371**, eaba8310 (2021).
- Lyu, M. et al. ILC3s select microbiota-specific regulatory T cells to establish tolerance in the gut. *Nature* **610**, 744–751 (2022).
- Jarade, A. et al. Inflammation triggers ILC3 patrolling of the intestinal barrier. *Nat. Immunol.* **23**, 1317–1323 (2022).
- Ou, R. & Murphy, K. M. What's in a name: clarifying the identity of ROR γ t⁺ antigen-presenting cells. *J. Exp. Med.* **222**, e20250760 (2025).
- von Moltke, J., Ji, M., Liang, H.-E. & Locksley, R. M. Tuft-cell-derived IL-25 regulates an intestinal ILC2–epithelial response circuit. *Nature* **529**, 221–225 (2016).
- Gerbe, F. et al. Intestinal epithelial tuft cells initiate type 2 mucosal immunity to helminth parasites. *Nature* **529**, 226–230 (2016).
- Howitt, M. R. et al. Tuft cells, taste-chemosensory cells, orchestrate parasite type 2 immunity in the gut. *Science* **351**, 1329–1333 (2016).
- Schneider, C. et al. A metabolite-triggered tuft cell–ILC2 circuit drives small intestinal remodeling. *Cell* **174**, 271–284 (2018).
- Van Dyken, S. J. et al. A tissue checkpoint regulates type 2 immunity. *Nat. Immunol.* **17**, 1381–1387 (2016).
- Vannella, K. M. et al. Combinatorial targeting of TSLP, IL-25, and IL-33 in type 2 cytokine-driven inflammation and fibrosis. *Sci. Transl. Med.* **8**, 337ra65 (2016).
- Taylor, B. C. et al. TSLP regulates intestinal immunity and inflammation in mouse models of helminth infection and colitis. *J. Exp. Med.* **206**, 655–667 (2009).
- Haber, A. L. et al. A single-cell survey of the small intestinal epithelium. *Nature* **551**, 333–339 (2017).
- Li, M. et al. Topical vitamin D3 and low-calcemic analogs induce thymic stromal lymphopoietin in mouse keratinocytes and trigger an atopic dermatitis. *Proc. Natl Acad. Sci. USA* **103**, 11736–11741 (2006).
- Shoshkes-Carmel, M. et al. Subepithelial telocytes are an important source of Wnts that supports intestinal crypts. *Nature* **557**, 242–246 (2018).
- McCarthy, N. et al. Distinct mesenchymal cell populations generate the essential intestinal BMP signaling gradient. *Cell Stem Cell* **26**, 391–402.e5 (2020).
- Bahar Halpern, K. et al. Lgr5⁺ telocytes are a signaling source at the intestinal villus tip. *Nat. Commun.* **11**, 1936 (2020).
- Melissari, M.-T. et al. Col6a1⁺CD201⁺ mesenchymal cells regulate intestinal morphogenesis and homeostasis. *Cell. Mol. Life Sci.* **79**, 1 (2021).
- Nussbaum, J. C. et al. Type 2 innate lymphoid cells control eosinophil homeostasis. *Nature* **502**, 245–248 (2013).
- Ricardo-Gonzalez, R. R. et al. Tissue-specific pathways extrude activated ILC2s to disseminate type 2 immunity. *J. Exp. Med.* **217**, e20191172 (2020).
- Buechler, M. B. et al. Cross-tissue organization of the fibroblast lineage. *Nature* **593**, 575–579 (2021).
- Gehart, H. et al. Identification of enteroendocrine regulators by real-time single-cell differentiation mapping. *Cell* **176**, 1158–1173 (2019).
- Bai, L. et al. Enteroendocrine cell types that drive food reward and aversion. *eLife* **11**, e74964 (2022).
- Smillie, C. S. et al. Intra- and inter-cellular rewiring of the human colon during ulcerative colitis. *Cell* **178**, 714–730 (2019).
- Korsunsky, I. et al. Cross-tissue, single-cell stromal atlas identifies shared pathological fibroblast phenotypes in four chronic inflammatory diseases. *Med* **3**, 481–518 (2022).
- Brubaker, P. L. Glucagon-like peptide-2 and the regulation of intestinal growth and function. *Compr. Physiol.* **8**, 1185–1210 (2018).
- O'Rahilly, S. The islet's bridesmaid becomes the bride: proglucagon-derived peptides deliver transformative therapies. *Cell* **184**, 1945–1948 (2021).
- Drucker, D. J. Mechanisms of action and therapeutic application of glucagon-like peptide-1. *Cell Metab.* **27**, 740–756 (2018).
- de Heuvel, E., Wallace, L., Sharkey, K. A. & Sigalet, D. L. Glucagon-like peptide 2 induces vasoactive intestinal polypeptide expression in enteric neurons via phosphatidylinositol 3-kinase-g signaling. *Am. J. Physiol. Endocrinol. Metab.* **303**, E9941005 (2012).
- Yusta, B. et al. Identification of glucagon-like peptide-2 (GLP-2)-activated signaling pathways in baby hamster kidney fibroblasts expressing the rat GLP-2 receptor. *J. Biol. Chem.* **274**, 30459–30467 (1999).
- Koehler, J. A., Harper, W., Barnard, M., Yusta, B. & Drucker, D. J. Glucagon-like peptide-2 does not modify the growth or survival of murine or human intestinal tumor cells. *Cancer Res.* **68**, 7897–7904 (2008).
- Futamura, K. et al. beta2-Adrenoceptor agonists enhance cytokine-induced release of thymic stromal lymphopoietin by lung tissue cells. *Int. Arch. Allergy Immunol.* **152**, 353–361 (2010).
- Ma, L., Zhen, J. & Sorisky, A. Regulators of thymic stromal lymphopoietin production by human adipocytes. *Cytokine* **136**, 155284 (2020).

34. Kondo, A. & Kaestner, K. H. Emerging diverse roles of telocytes. *Development* **146**, dev175018 (2019).
35. Beumer, J. et al. BMP gradient along the intestinal villus axis controls zonated enterocyte and goblet cell states. *Cell Rep.* **38**, 110438 (2022).
36. Krausgruber, T. et al. Structural cells are key regulators of organ-specific immune responses. *Nature* **583**, 296–302 (2020).
37. Gribble, F. M. & Reimann, F. Enteroendocrine cells: chemosensors in the intestinal epithelium. *Annu. Rev. Physiol.* **78**, 277–299 (2016).
38. Beumer, J. et al. Enteroendocrine cells switch hormone expression along the crypt-to-villus BMP signalling gradient. *Nat. Cell Biol.* **20**, 909–916 (2018).
39. Leen, J. L. S. et al. Mechanism of action of glucagon-like peptide-2 to increase IGF-I mRNA in intestinal subepithelial fibroblasts. *Endocrinology* **152**, 436–446 (2011).
40. Wismann, P. et al. The endogenous preproglucagon system is not essential for gut growth homeostasis in mice. *Mol. Metab.* **6**, 681–692 (2017).
41. Shin, E. D., Estall, J. L., Izzo, A., Drucker, D. J. & Brubaker, P. L. Mucosal adaptation to enteral nutrients is dependent on the physiologic actions of glucagon-like peptide-2 in mice. *Gastroenterology* **128**, 1340–1353 (2005).
42. Nelson, D. W. et al. Insulin-like growth factor I and glucagon-like peptide-2 responses to fasting followed by controlled or ad libitum refeeding in rats. *Am. J. Physiol. Regul. Integr. Comp. Physiol.* **294**, R1175–R1184 (2008).
43. Tsai, C. H., Hill, M., Asa, S. L., Brubaker, P. L. & Drucker, D. J. Intestinal growth-promoting properties of glucagon-like peptide-2 in mice. *Am. J. Physiol.* **273**, E77–E84 (1997).
44. Rankin, L. C. & Artis, D. Beyond host defense: emerging functions of the immune system in regulating complex tissue physiology. *Cell* **173**, 554–567 (2018).
45. Schneider, C., O’Leary, C. E. & Locksley, R. M. Regulation of immune responses by tuft cells. *Nat. Rev. Immunol.* **19**, 584–593 (2019).
46. Florsheim, E. B., Sullivan, Z. A., Khoury-Hanold, W. & Medzhitov, R. Food allergy as a biological food quality control system. *Cell* **184**, 1440–1454 (2021).
47. Nadjisombati, M. S. et al. Detection of succinate by intestinal tuft cells triggers a type 2 innate immune circuit. *Immunity* **49**, 33–41 (2018).
48. Luo, X.-C. et al. Infection by the parasitic helminth *Trichinella spiralis* activates a Tas2r-mediated signaling pathway in intestinal tuft cells. *Proc. Natl Acad. Sci. USA* **116**, 5564–5569 (2019).
49. Chudnovskiy, A. et al. Host–protozoan interactions protect from mucosal infections through activation of the inflammasome. *Cell* **167**, 444–456 (2016).
50. Ramanan, D. et al. Helminth infection promotes colonization resistance via type 2 immunity. *Science* **352**, 608–612 (2016).

Publisher’s note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2025

Methods

Mice

C57BL/6J mice were purchased from the Jackson laboratory (JAX, catalog number 000664) or mutant alleles backcrossed more than eight generations to C57BL/6J. Red5 or B6(C)-*Il5^{tm1.1(cre)Lky}/J*¹⁹, Smart13 or B6.129S4(C)-*Il13^{tm2.1Lky}/J*¹⁰, *Tslpr^{-/-}* (ref. 51), backcrossed *Il4ra^{-/-}* or BALB/c-*Il4ra^{tm1Sz}/J* (JAX, catalog number 003514)⁶, and *Il3ra1^{-/-}* (ref. 52 mice generated or obtained by this laboratory have been described. *Vil1^{Cre}* or B6.Cg-Tg(*Vil1-cre*)997Gum/J (JAX, catalog number 004586), *Pdgfra^{CreERT2}* or B6.129S-*Pdgfra^{tm1.1(cre/ERT2)Blh}/J* (JAX, catalog number 032770), *Nes^{CreERT2}* or C57BL/6-Tg(*Nes-cre/ERT2*)KEisc/J (JAX, catalog number 016261) were purchased from the Jackson laboratory. C57BL/6 *Pou2f3^{-/-}* (rederived from C57BL/6NTac *Pou2f3^{tm1.1(KOMP)Vicg}*) mice⁹ were provided by M. S. Anderson. C57BL/6 *Tslpr^{fl/fl}* mice⁵³ were provided by S. F. Ziegler. C57BL/6 *Glp2r^{-/-}* mice⁵⁴ were provided by D. Drucker. *Lgr5-eGFP* mice⁵⁵ were provided from Genentech by F. de Sauvage and O. D. Klein. Triple-mutant *Vil1^{flp} Cck^{Cre}* (JAX, catalog number 012706) *R26^{Dual-hM3Dq}* (JAX, catalog number 026942) mice²³ were provided by Z. A. Knight.

Flare-TSLP mice were generated by homologous gene targeting in C57BL/6 embryonic stem cells based on a reporter cassette used to target the *Il25* locus⁶. Briefly, a 2.34-kb 3' homologous arm in the 3' untranslated region of *Tslp* was amplified from C57BL/6 genomic DNA and a 3.79-kb fragment from exon 1 to the end of coding sequence of exon 5 was amplified to serve as the 5' homologous arm. A 5' *loxP* site (target site for Cre recombinase) was inserted 187 bp upstream of exon 4 within the modified 5' arm. The reporter cassette encoded (5' to 3') a *loxP* site (3' *loxP* in final construct), encephalomyocarditis virus internal ribosomal entry site (IRES) element, tandem RFP or tdTomato, bovine growth hormone poly(A) signal and an frt-flanked neomycin selection cassette, and was subcloned between the modified 5' and 3' homologous arms

with correct orientation using the multiple cloning site of a basal targeting vector, pKO915-DT (Lexicon Genetics). After linearization by *NotI*, the reporter cassette was electroporated into C57BL/6 embryonic stem cells. Following growth on irradiated mouse embryonic fibroblast feeders and selection in G418-containing medium, neomycin-resistant clones were screened for correct 5' and 3' homologous recombination by long-range PCR and further confirmed for retention of the 5' *loxP* site. One clone was injected into albino C57BL/6 blastocysts to generate chimeras and males with coats with the highest black-to-white coat color ratio were bred with B6.129S4-*Gt(Rosa)26Sor^{tm1(FLP1)Dym}/RainJ* females (JAX, catalog number 009086) to excise the neomycin resistance cassette. Offspring with germline transmission and confirmed neo⁻-deleted *Flare-Tslp* allele were backcrossed to C57BL/6 mice to cross out the FLP1 allele. *Flare-Tslp* was genotyped using primers 5' *loxP fw*: 5'-GGAACGAAGTTGAAACACCACGACC-3' and 5' *loxP rev*: 5'-GGGGATGGGGATAGGAGGAAAGAC-3', yielding a 277-bp mutant or a 243-bp WT band. After Cre-*loxP*-mediated deletion, the floxed-out *Tslp* allele can be detected by PCR using primers 5' *loxP fw* (as above) and *IRES rev*: 5'-TTGTTGAATACGCTTGAGGAGA GCC-3', yielding a 596 base pair band, whereas a nonrecombined floxed band is 2,878 base pairs.

Mice were maintained in the University of California, San Francisco (UCSF) specific pathogen-free animal facility in accordance with the guidelines established by the Institutional Animal Care and Use Committee and Laboratory Animal Resource Center. Unless otherwise specified, mice were kept on 12-h light–dark cycles and ad libitum water and chow (PicoLab, catalog number 5053). The facility was maintained at temperature 68–79 °F and 30–70% humidity. Pooled experiments included female and male mice at 8–12 weeks old. All experimental procedures were approved by the Laboratory Animal Resource Center at the UCSF.

Antibodies

Antibodies	Supplier	Catalog number	Dilution
Brilliant violet 421 anti-mouse CD19 antibody	BioLegend	115549	1:200
Brilliant violet 421 anti-mouse TER-119/erythroid cells antibody	BioLegend	116234	1:200
Brilliant violet 421 anti-mouse Ly-6G/Ly-6C (Gr-1) antibody	BioLegend	108445	1:200
Brilliant violet 605 anti-human/mouse/rat CD278 (ICOS) antibody	BioLegend	313538	1:100
Brilliant violet 421 anti-mouse/human CD11b antibody	BioLegend	101251	1:200
Pacific blue anti-mouse CD11c antibody	BioLegend	117322	1:200
Pacific blue anti-mouse CD49b (pan-NK cells) antibody	BioLegend	108918	1:200
Brilliant violet 421 anti-mouse CD335 (Nkp46) antibody	BioLegend	137612	1:200
Brilliant violet 421 anti-mouse NK1.1 antibody	BioLegend	108741	1:200
Brilliant violet 421 anti-mouse TCR γ/δ antibody	BioLegend	118120	1:200
Pacific blue anti-mouse FcεR1α antibody	BioLegend	134314	1:200
Pacific blue anti-mouse F4/80 antibody	BioLegend	123124	1:200
BUV395 rat anti-mouse CD45	BD	564279	1:200
PE/Cyanine7 anti-mouse/human KLRG1 (MAFA) antibody	BioLegend	138416	1:100
APC anti-human CD4 antibody	BioLegend	300514	1:50
Ki-67 monoclonal antibody (SolA15), FITC, eBioscience	Invitrogen	11-5698-82	1:1000
APC anti-mouse CD201 (EPCR) antibody	BioLegend	141506	1:200
PerCP/cyanine5.5 anti-mouse CD31 antibody	BioLegend	102420	1:200
Brilliant violet 421 anti-mouse CD31 antibody	BioLegend	102424	1:200
Brilliant violet 605 anti-mouse CD140α antibody	BioLegend	135916	1:100
Alexa Fluor 488 anti-mouse LYVE1 antibody	eBiosciences	53-0443-82	1:100
Alexa Fluor 488 anti-mouse podoplanin antibody	BioLegend	127406	1:100
BV786 rat anti-mouse CD34	BD	742971	1:100

Antibodies	Supplier	Catalog number	Dilution
Alexa Fluor 647 anti-mouse CD326 (EPCAM) antibody	BioLegend	118212	1:100
Alexa Fluor 488 anti-mouse CD326 (EPCAM) antibody	BioLegend	118210	1:100
Alexa Fluor 488 Rat IgG2a, κ isotype Ctrl antibody	BioLegend	400525	1:100
Anti-DCAMKL1 (DCLK1) antibody	Abcam	ab31704	1:500
Mouse podoplanin antibody	R&D	AF3244	1:250
Mouse PDGFR alpha antibody	R&D	MAB1062	1:250
Mouse EPCR (CD201) antibody	R&D	AF2749	1:200
CD34 monoclonal antibody (RAM34), eBioscience	ThermoFisher	14-0341-82	1:500
Anti-green fluorescent protein antibody	Aveslabs	GFP-1020	1:500
Living colors DsRed polyclonal antibody	TakaraBio	632496	1:250
mCherry monoclonal antibody (16D7)	ThermoFisher	M11217	1:500
TSLP monoclonal antibody	Invitrogen	MA5-23779	1:100

TSLP antibody labeling

Mouse TSLP antibody for flow cytometric analysis was labeled using Alexa Fluor 488 Antibody Labeling Kit (ThermoFisher, catalog number A20181) following the manufacturer’s protocol.

Flow cytometry and FACS

Small intestinal tissues were harvested at indicated times and single-cell suspensions of epithelium or lamina propria were obtained. In brief, 6-cm lengths of proximal jejunum or distal ileum were collected and flushed with ice-cold PBS. After removing the Peyer’s patches, gut tissues were opened and washed with PBS to remove lumen content and mucus. Segments were incubated while rocking for 15 min at 37 °C in 20 ml HBSS (Ca²⁺-free and Mg²⁺-free) containing 5% FCS, 10 mM HEPES (Sigma, catalog number H3537), 10 mM EDTA (Corning, catalog number 46-034-CI) and 10 mM dithiothreitol (DTT; Sigma, catalog number D0632). Tissue segments were vortexed vigorously for 30 s and supernatants collected and centrifuged. The pelleted epithelial fraction was washed once with PBS before staining for flow cytometry. The remaining tissue segments were incubated a second time while rocking for 15 min at 37 °C in 10 ml of the same HBSS/FBS/EDTA/DTT medium and then vortexed for 30 s. The tissue segments were transferred to new tubes and incubated while rocking for 5 min at 37 °C in 20 ml HBSS (with Ca²⁺ and Mg²⁺) containing 2% FCS and 10 mM HEPES before transferring to new tubes containing 5 ml HBSS (with Ca²⁺ and Mg²⁺), 3% FCS, 10 mM HEPES, 30 μ g ml⁻¹ DNase (Roche, catalog number 10104159001) and 0.1 U ml⁻¹ Liberase TM (Roche, catalog number 5401127001) in C tubes (Miltenyi Biotec, catalog number 130-096-334). Samples were processed with gentleMax Octo Dissociator program ‘intestine.’ Lamina propria cells were pelleted, washed and filtered with 70 μ m cell strainers before staining for flow cytometry. Standard surface and/or intracellular staining protocols were used followed by 4’,6-diamidino-2-phenylindole (DAPI) or Live/Dead Viability dye (ThermoFisher) staining for dead cell exclusion. Samples were processed using a BD LSRFortessa (Bectin Dickinson) flow cytometer and analyzed using FlowJo software. Cell sorting experiments were performed on MoFlo (Beckman Coulter) or BD FACSAria.

Primary mouse fibroblast culture

sILP cells were isolated as above. Cells were suspended in fibroblast culture media (DEME/F12 supplement with 10% FBS and penicillin/streptomycin) and incubated at 37 °C with 5% CO₂ for 4 h to allow fibroblast adherence before washing off nonadherent cells. The remaining adherent cells were incubated in fresh culture medium. Cells were plated at 1 \times 10⁵ cells per well in 48-well plates or 2 \times 10⁵ cells per well in 24-well plates and incubated overnight before use.

Human intestinal fibroblast cell lines

Study approval. The study was conducted in accordance with the principles of the Declaration of Helsinki and was approved by the Institutional Review Board of the University of California, San Francisco (19-27302). All participants provided written informed consent before inclusion.

Study participants and biospecimen collection

Patients underwent colonoscopy or sigmoidoscopy for noninflammatory indications (for example, colorectal cancer screening) and were referred as healthy controls (HC). Baseline demographic information for the study participants are provided in Supplementary Table 1. Patients consented to publish deidentified patient demographics including age at the time of sample collection, sex, diagnosis and medical center. Demographic options were defined by the investigators and participants chose their classifications.

Sample collection and storage

Biopsies from colon and ileum were obtained with standard cold endoscopic biopsy forceps and collected in a conical tube with Basal Media (Advanced DMEM/F12 with nonessential amino acids and sodium pyruvate (Thermo, catalog number 12634-010), 2 mM Glutamax (Thermo, catalog number 35050061), 10 mM HEPES (Corning), penicillin-streptomycin-neomycin (PSN) Antibiotic Mixture (Thermo, catalog number 15640055), 100 μ g ml⁻¹ Normocin (Invivogen, catalog number ant-nr-2), 1 mM N-acetylcysteine (Sigma-Aldrich, catalog number A9165)) with 10 μ M Y-27632 (MedChem Express) at 4 °C. Samples were immediately placed on ice and transported to the laboratory for processing as described⁵⁶. Biopsies were transferred into cryovials containing freezing medium (90% (v/v) FCS, 10% (v/v) dimethylsulfoxide (DMSO) and 10 μ M Y-27632) and placed immediately into a freezing container (Coolcell) and stored at -70 °C for up to 4 weeks before transferring to liquid nitrogen cryostorage until further processing.

Establishment of patient-derived intestinal fibroblast cell lines

Biopsies were thawed for 2.5 min with gentle agitation in a 37 °C water bath, transferred to a 50-ml tube, washed twice with basal medium containing 10 μ M Y-27632, and then incubated in 2 ml digestion buffer (basal medium with 10 μ M Y-27632, 600 U ml⁻¹ Collagenase IV (Worthington, catalog number LS004189), 0.1 mg ml⁻¹ DNase I (Sigma-Aldrich, catalog number D4513)). Biopsies were digested for 20 min at 37 °C in a shaking incubator set at 225 rpm. Samples were pipetted vigorously after incubation. The suspension was passed through a 100- μ m strainer (pluriSelect, catalog number 43-100100-40) over a 5-ml tube, and the filter was washed twice with HBSS (Corning),

containing 0.1 mg ml⁻¹ DNase I. The suspension was centrifuged at 450g for 5 min at 4 °C. One additional wash was performed in HBSS containing 0.1 mg ml⁻¹ DNase I and centrifuged at 450g for 5 min at 4 °C. The pellet was resuspended in prewarmed human fibroblast culture medium (EMEM-Eagle's minimal essential medium with L-glutamine (Quality Biological, catalog number 112-018-101), PSN, 100 µg ml⁻¹ Normocin (InvivoGen, catalog number ant-nr-2), FCS 20% (v/v) and 50 ng ml⁻¹ recombinant human FGF-basic (PeproTech, catalog number 100-18B)) and plated in a 24-well plate. The culture medium was replaced after 24 h in culture. After expansion, fibroblast lines were transferred to T25 flasks and passaged twice a week at a ratio of 1:3. Aliquots of low-passage (<8) patient-derived intestinal fibroblasts were cryopreserved in 90% FCS with 10% DMSO.

Cells used in this paper were thawed from frozen stocks, expanded in human fibroblast culture medium and maintained in cell incubator at 37 °C with 5% CO₂. Cells were seeded at 2 × 10⁵ cells per well in 24-well plates and incubated overnight before use.

TSLP ELISA from mouse intestinal homogenates and explants

Samples from proximal jejunum (2 cm) or distal ileum (3 cm) were weighed and harvested in RIPA (ThermoFisher, catalog number 89900) buffer containing proteinase inhibitors in M tubes (Miltenyi Biotec, catalog number 130-096-335) followed by dissociation with gentleMax Octo Dissociator program 'protein 1'. Supernatants from the homogenates were collected following brief centrifugation and stored at -80 °C. Intestinal explants were cultured in 96-well plates with complete RPMI 1640 medium supplemented with 10% FBS for 5 h at 37 °C in a 5% CO₂ incubator. After incubation, supernatants were collected following brief centrifugation and stored at -80 °C. TSLP ELISA was performed (BioLegend Deluxe TSLP ELISA kit; BioLegend, catalog number 434104) with undiluted homogenates or tissue explant culture supernatants. Protein concentrations were normalized by tissue weight of the starting samples.

Drug treatments

MC903 (Calcipotriol, Tocris Biosciences, catalog number 2700; 100 µM working concentration in ethanol) was applied topically on shaved back skin using 60 µl per mouse daily for 7 consecutive days¹⁴. Stabilized human GLP-2[Gly2] (Teduglutide, Echelon Bioscience, catalog number 471-21) was given at 10 µg per mouse subcutaneously (s.c.) in 100 µl DMSO-PBS solution consecutively for 3 days or as indicated at 25 µg per mouse intraperitoneally (i.p.) 2 h before collecting tissues. Tamoxifen (Sigma, catalog number T5648) was administered at 3 mg per mouse or 100 mg kg⁻¹ in 100 µl corn oil i.p. daily for 7 days; mice were rested for 3–5 days before experimental use. Alternatively, tamoxifen diet (Inotive, catalog number TD.130858) was given to mice ad libitum for at least 4 weeks before experimental use. CNO (Cayman, catalog number 16882) was given 5 mg kg⁻¹ i.p. for 3 consecutive days. EdU (Sigma, catalog number 900584) was given i.p. at 1 mg per mouse in 100 µl PBS 24 h before tissue harvests.

For signaling pathway analysis, mouse fibroblasts, mouse intestinal explants and human fibroblasts were treated with agonists including GLP-2[Gly2] 500 nM and forskolin 100 µM (Tocris, catalog number 1099), or inhibitors including 666-15 10 µM (Tocris, catalog number 5661), LY294002 75 µM (Tocris, catalog number 1130), rapamycin 2.5 µM (Biogems, catalog number 1672186) and PD98059 50 µM (TCI, catalog number R0097) in 200 µl of culture medium. Inhibitors were added 1 h before agonists. For TSLP measurement by ELISA (BioLegend, catalog number 521904 for human TSLP; BioLegend, catalog number 434104 for mouse TSLP), supernatants were collected 5 h after agonist stimulation. For TSLP measurement by intracellular staining and flow cytometric analysis, cells were incubated for 4 h with Brefeldin A (BioLegend, catalog number 42601) given 1 h after agonists before cells were harvested and analyzed using standard cell surface and intracellular staining.

N. brasiliensis and *T. muris* infection

N. brasiliensis was maintained in rats and purified as described⁶. Mice were injected s.c. with 500 purified L3 larvae and small intestines were collected on days 5, 7 and 12 after infection for analysis of tuft cells as described⁶.

Two hundred embryonated *T. muris* eggs were orally gavaged per mouse. Cecum and large intestines were analyzed 21 days later.

Fasting-feeding protocols

For measuring small intestinal TSLP, mice were fasted 16 h overnight before oral gavage with 500 µl food slurry (modified powdered diet formula TekladTD.88232 with comparable nutrient distribution as standard chow diet) in water containing 1.965 kcal for ~1% daily caloric intake) or water as a volumetric control. For small intestinal ILC2 activation, mice were fasted overnight for 16 h before restoring access to standard chow diet (PicoLab, catalog number 5058) and water ad libitum or maintaining on water. Tissues were harvested at indicated timepoints. Overnight fasting followed by water gavage or water access ad libitum was designated the 'Fasted' control group; overnight fasting followed by food gavage or food access ad libitum was designated the 'Fasted+refed' group.

Tissue immunofluorescence

Tissues were fixed in 4% paraformaldehyde (PFA; Electron, catalog number 15710) for 2 h at room temperature or longer at 4 °C, washed three times in PBS and incubated in 30% (w/v) sucrose-PBS solution overnight at 4 °C. Tissues were embedded in optimal cutting temperature compound (OCT; VWR, catalog number 25608-930) on dry ice and stored at -80 °C. Skin samples were cut into 1 cm × 2 cm pieces before fixation. Samples from the small intestine (~10 cm) or proximal large intestine were cut and flushed with PBS before fixation and coiled into 'Swiss rolls' before embedding. OCT-embedded tissues were cut into 8–14-µm regular sections or 100-µm thick sections using a Cryostat (Leica, catalog number CM3050 S) and stored at -80 °C. Sections were incubated with 0.02% Triton X-100 (Sigma, catalog number X100) in TNB buffer (Tris-NaCl blocking buffer: 0.1 M Tris-HCl, pH 7.5; 0.15 M NaCl; blocking reagent, Akoya Biosciences, catalog number FP1020) for permeabilization followed by blocking with 5% serum matching the source of the secondary antibodies for 30 min at room temperature. After a 2-h incubation with primary antibodies at room temperature, sections were washed three times in PBS and incubated with secondary antibodies for 1 h at room temperature. Sections were incubated with DAPI for 5 min and washed three times in PBS before mounting. EdU staining was performed using Click-IT Plus EdU imaging kit (Invitrogen, catalog number C10637) following manufacture protocol. Stained samples were visualized using a Nikon A1R confocal microscope with a ×20 objective and analyzed using NIS Element and ImageJ software. Thick section staining protocols were as described⁵⁷ and imaging was processed using Imaris software three-dimensional analysis workstation for Cell Biology Research (Oxford Instruments).

Tuft cell quantifications

For tuft cell quantification, images were acquired using the ×20 objective and total doublecortin-like kinase 1 (DCLK1)⁺EPCAM⁺ cells in each field were counted and normalized by total number of villus-crypt axes in the field.

qPCR and gene expression analysis

For analysis of designated populations, cells were purified by FACS directly into lysis buffer followed by RNA extraction using a Qiagen kit (catalog number 74034). cDNA was synthesized using Invitrogen Super-Script IV VILO kit (Invitrogen, catalog number 11754250). qPCR reactions were performed using Taqman probes (Invitrogen) and Taqman gene expression master mix (Invitrogen, catalog number 4369016). Transcripts were normalized to 18S expression. Probe information is as follows:

Gene	Probe
<i>Rspo3</i>	Mm00661105_m1
<i>Lgr5</i>	Mm00438890_m1
<i>Foxl1</i>	Mm00514937_s1
<i>Bmp4</i>	Mm00432087_m1
<i>Wnt2b</i>	Mm00437330_m1
<i>Wnt5a</i>	Mm00437347_m1
<i>Grem1</i>	Mm00488615_s1
<i>Bmp7</i>	Mm00432102_m1
18S	Mm03928990_g1
<i>Tslp</i>	Mm00498739_m1
Nes	Mm00450205_m1

scRNA-seq and cell hashing

Live tdTomato⁺ cells were purified using FACS and labeled with unique barcoded H-2/CD45 TotalSeq-B anti-mouse hashtag antibodies (Biolegend catalog numbers 155831, 155833, 155835, 155837, 155839, 155841) for multiplexing. The cells were processed according to the manufacturer’s instructions using the Chromium Next Gel Beads-in-Emulsion (GEM) Single Cell 3’ Kit v.3.1 (10x Genomics, catalog number 1000269). Briefly, cell counts were performed using a hemocytometer and the desired number of cells was injected into microfluidic chips (10x Genomics, catalog number 1000127) to create GEMs with the 10x Chromium controller. RT was performed on the GEMs followed by purification and amplification of the RT products. Cell multiplexing oligo (CMO) DNA was separated from cDNA using size selection with SPRI select beads (Beckman Coulter, catalog number B23318). Gene expression libraries and CMO libraries were generated separately and profiled using the Bioanalyzer High Sensitivity DNA kit (Agilent Technologies, catalog number 5067-4626). Before sequencing, the gene expression and corresponding CMO libraries were mixed in a 4:1 ratio. Sequencing of the mixed libraries was conducted on the Illumina NovaSeq X platform.

scRNA-seq data analysis

Sample demultiplexing, alignments and unique molecular identifier counts were performed using the 10x Genomics Cell Ranger pipeline (v.8.1.0) based on the latest mouse genome (mm39) with a custom reference to include the tdTomato sequence. Doublets were removed using the DoubletFinder (v.2.0.4) algorithm and ambient RNA was eliminated with the SoupX algorithm (v.1.6.2) using default parameters. After preprocessing datasets were analyzed using the Seurat single-cell analysis pipeline (v.5.0.3) in R (v.4.3.1). Quality control involved excluding low-quality cells based on the following criteria: nFeature_RNA between 1,000 and 7,500, nCount_RNA between 2,500 and 40,000 and mitochondrial contamination below 10%. Following filtering, data were normalized using the NormalizeData function with the ‘LogNormalize’ method and the 2,000 most variable genes were identified and scaled before performing principal component analysis. Thirty principal components were used for graph-based clustering (resolution = 0.5) followed by uniform manifold approximation and projection dimensionality reduction.

An additional filtering step based on *tdTomato* expression removed contaminant clusters with fewer than 0.01% *tdTomato*⁺ cells and fewer than five absolute *tdTomato*⁺ cells. Initial cell type annotation was conducted automatically using scType³⁸ followed by manual refinement. Differential gene expression (DE) analysis between cell types in the small intestine samples were conducted using Wilcoxon rank sum tests as implemented in the Seurat function FindAllMarkers. For DE analysis between tissues, counts from each sample were summed and followed by pseudobulk DE analysis using limma-voom.

Public scRNA-seq data analysis

Mouse^{15,16} and human^{24,25} datasets (raw or preprocessed data) were obtained from the single-cell portal of the Broad Institute (<https://singlecell.broadinstitute.org>). Data were reanalyzed and plotted using Seurat package in the R program.

Statistics

Quantitative data were represented as mean ± s.e.m. of at least duplicate biological replicates. Student’s *t*-test was used for analyses of difference between two groups. Analysis of variance (ANOVA) was used for analyzing experiments with more than two groups adjusted for multiple comparisons. Significance level was set at α = 0.05. Statistical analysis was performed using Prism (GraphPad Software). **P* < 0.05, ***P* < 0.01, ****P* < 0.001, *****P* < 0.0001.

Reporting summary

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

Data availability

All raw and processed scRNA-seq data generated in this study have been deposited in the NCBI Gene Expression Omnibus (GEO) under accession number [GSE280635](https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE280635). All the other data supporting the findings of this study are available upon request.

Code availability

This study did not generate new algorithm or code. The custom code and scripts used for analyses and generating related graphs are available upon requests.

References

51. Carpino, N. et al. Absence of an essential role for thymic stromal lymphopoietin receptor in murine B-cell development. *Mol. Cell Biol.* **24**, 2584–2592 (2004).

52. Ricardo-Gonzalez, R. R. et al. Innate type 2 immunity controls hair follicle commensalism by *Demodex* mites. *Immunity* **55**, 1891–1908 (2022).

53. Han, H., Thelen, T. D., Comeau, M. R. & Ziegler, S. F. Thymic stromal lymphopoietin-mediated epicutaneous inflammation promotes acute diarrhea and anaphylaxis. *J. Clin. Invest.* **124**, 5442–5452 (2014).

54. Yusta, B. et al. ErbB signaling is required for the proliferative actions of GLP-2 in the murine gut. *Gastroenterology* **137**, 986–996 (2009).

55. Tian, H. et al. A reserve stem cell population in small intestine renders Lgr5-positive cells dispensable. *Nature* **478**, 255–259 (2011).

56. Mennillo, E. et al. Single-cell and spatial multi-omics highlight effects of anti-integrin therapy across cellular compartments in ulcerative colitis. *Nat. Commun.* **15**, 1493 (2024).

57. Li, W., Germain, R. N. & Gerner, M. Y. High-dimensional cell-level analysis of tissues with Ce3D multiplex volume imaging. *Nat. Protoc.* **14**, 1708–1733 (2019).

58. Ianevski, A., Giri, A. K. & Aittokallio, T. Fully-automated and ultra-fast cell-type identification using specific marker combinations from single-cell transcriptomic data. *Nat. Commun.* **13**, 1246 (2022).

59. Small intestine from C57BL/6 mice (v1, 150×150), Visium HD Spatial Gene Expression Library by Space Ranger v3.0. 10x Genomics <https://www.10xgenomics.com/datasets/visium-hd-cytassist-gene-expression-libraries-of-mouse-intestine?noTrack=true> (2024).

Acknowledgements

We thank A. Ma, A. B. Molofsky and D. J. Julius for review of the paper; A. B. Molofsky, M. S. Anderson, Z. A. Knight, S. F. Ziegler,

F. de Sauvage and O. D. Klein for mouse lines; J. F. Urban Jr and P. Loke for providing *Trichuris muris* embryonated eggs; current and former members of the Locksley laboratory for comments; UCSF core facilities including the Flow Cytometry Core, the Functional Genomic CoLab and the Center for Advanced Technology (CAT) for technical and bioinformatic support; the UC-Davis Bioinformatics Core for bioinformatic analyses; M. Conesco for mouse colony support; and G. W. Kim and A. Luong for administrative support. This work was supported by National Institutes of Health grant AIO26918 and HL107202 (R.M.L.), Howard Hughes Medical Institute (R.M.L.), Sandler Basic Asthma Research Center at University of California, San Francisco (R.M.L.), Canadian Institute Health Research (CIHR) grant 154321 (D.J.D.), Kenneth Rainin Foundation (M.G.K.) and Cancer Research Institute Fellowship (C.L.).

Author contributions

C.L. conceived the study, designed and performed experiments, analyzed and interpreted the data, and wrote the paper. H.-E.L. generated the Flare-TSLP reporter mice and participated in experiments. E.M. generated human intestinal fibroblast cell lines and wrote the corresponding methods. M.J. and Z.-E.W. participated in experiments. D.J.D. provided key reagents, analysis and editorial input. M.G.K. provided key reagents and editorial input. R.M.L. conceived and directed the studies with C.L., analyzed and interpreted data, and wrote the paper with C.L. with input from the co-authors.

Competing interests

D.J.D. is a party to a GLP-2 license agreement between Takeda Pharmaceuticals, the University Health Network and the University of Toronto. M.G.K. receives research support from Eli Lilly. M.G.K. has consulted for Sonoma Biotherapeutics, Morphic Therapeutic, Santa Ana Bio, Spyre, Cellarity, Vedanta and CytoMx, and he is on the scientific advisory board for Switchback Therapeutics. The other authors declare no competing interests.

Additional information

Extended data is available for this paper at <https://doi.org/10.1038/s41590-025-02328-y>.

Supplementary information The online version contains supplementary material available at <https://doi.org/10.1038/s41590-025-02328-y>.

Correspondence and requests for materials should be addressed to Richard M. Locksley.

Peer review information *Nature Immunology* thanks the anonymous reviewers for their contribution to the peer review of this work. Primary Handling Editor: S. Houston, in collaboration with the *Nature Immunology* team.

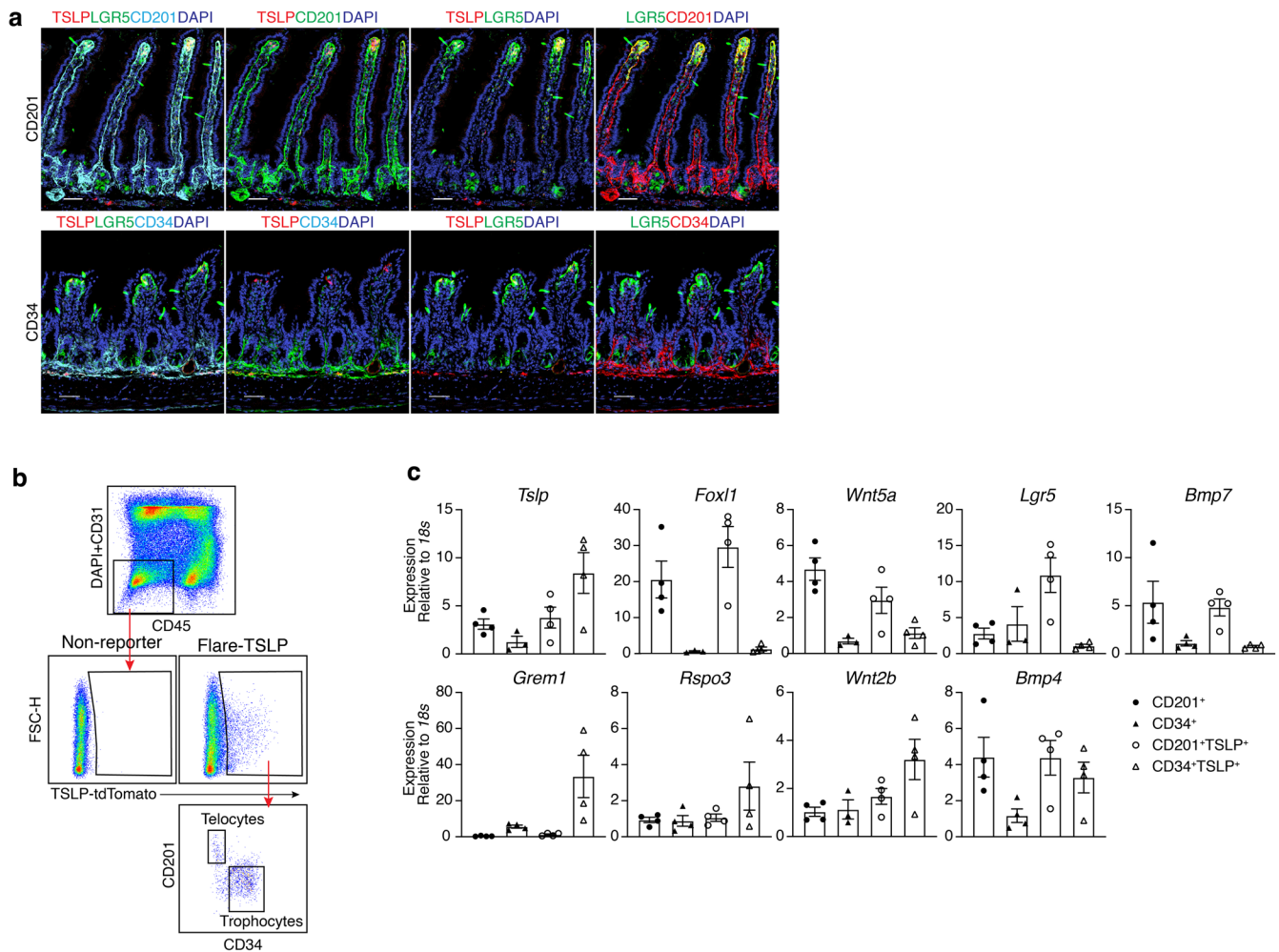
Reprints and permissions information is available at www.nature.com/reprints.



Extended Data Fig. 1 | Characterization of Flare-TSLP reporter mice.

a, Representative imaging showing jejunum (**top**) and ileum (**bottom**) of Flare-TSLP mice or non-reporter control mice. Red, TSLP-tdTomato; Green, PDPN; Blue, DAPI. 20x objective. Scale bar, 50 μ m. **Right panel**, max intensity projection of three-dimensional imaging showing jejunal tissues from Flare-TSLP mice; scale bar, 100 μ m. **b-d**, Flow cytometric analysis of CD45⁺ and stromal populations in jejunal and ileal lamina propria in Flare-TSLP or non-reporter control (ctrl) mice. **b**, CD45⁺ cells were CD31⁺ endothelial cells. CD45⁺CD31⁻ stromal cells were PDGFRa⁺cKit⁻ (non-Cajal cells, which are cKit⁺). **c**, Proportions of CD45⁺CD31⁺, CD45⁺CD31⁻, CD45⁻CD31⁺, and CD45⁻CD31⁻ cells in TSLP-tdTomato⁺ cells represented by pie charts. **d**, TSLP-tdTomato⁺CD45⁺CD31⁺ cells were PDPN⁺ and LYVE1⁺ consistent with lymphatic endothelial cells (LECs).

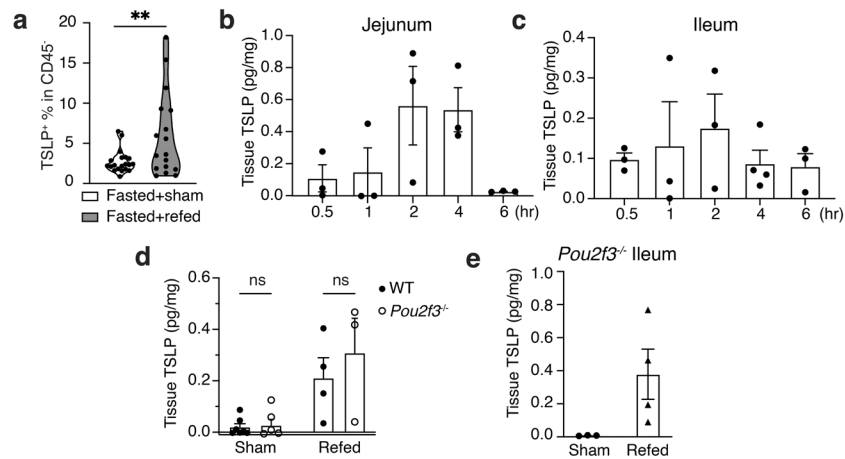
TSLP-tdTomato⁺CD45⁺ cells are CD31⁺FSC^{hi} large cells consistent with a non-hematopoietic lineage. **e**, Representative imaging showing back skin of Flare-TSLP mice treated with ethanol or calcipotriol (MC903) for 7 days. Red, TSLP-tdTomato; Blue, DAPI. 20x objective. Scale bar, 100 μ m. **f**, Flow cytometric analysis of epithelial TSLP expression in proximal small intestine in naive or on day 7 and 12 post *N. brasiliensis* infection in Flare-TSLP or non-reporter mice. d.p.i., day post infection. **g-h**, Representative imaging of jejunum in Flare-TSLP or non-reporter mice on day 5 after *N. brasiliensis* (**g**) and in cecum day 21 after *T. muris* infection (**h**). Red, TSLP-tdTomato; Green, PDGFRa; White, EPCAM; Blue, DAPI. 20x objective. Scale bar, 100 μ m. Inset scale bar, 50 μ m. All data are biological replicates, n \geq 3. Data are representative of at least two independent experiments.



Extended Data Fig. 2 | Analysis of CD201⁺ telocytes and CD34⁺ trophocytes.

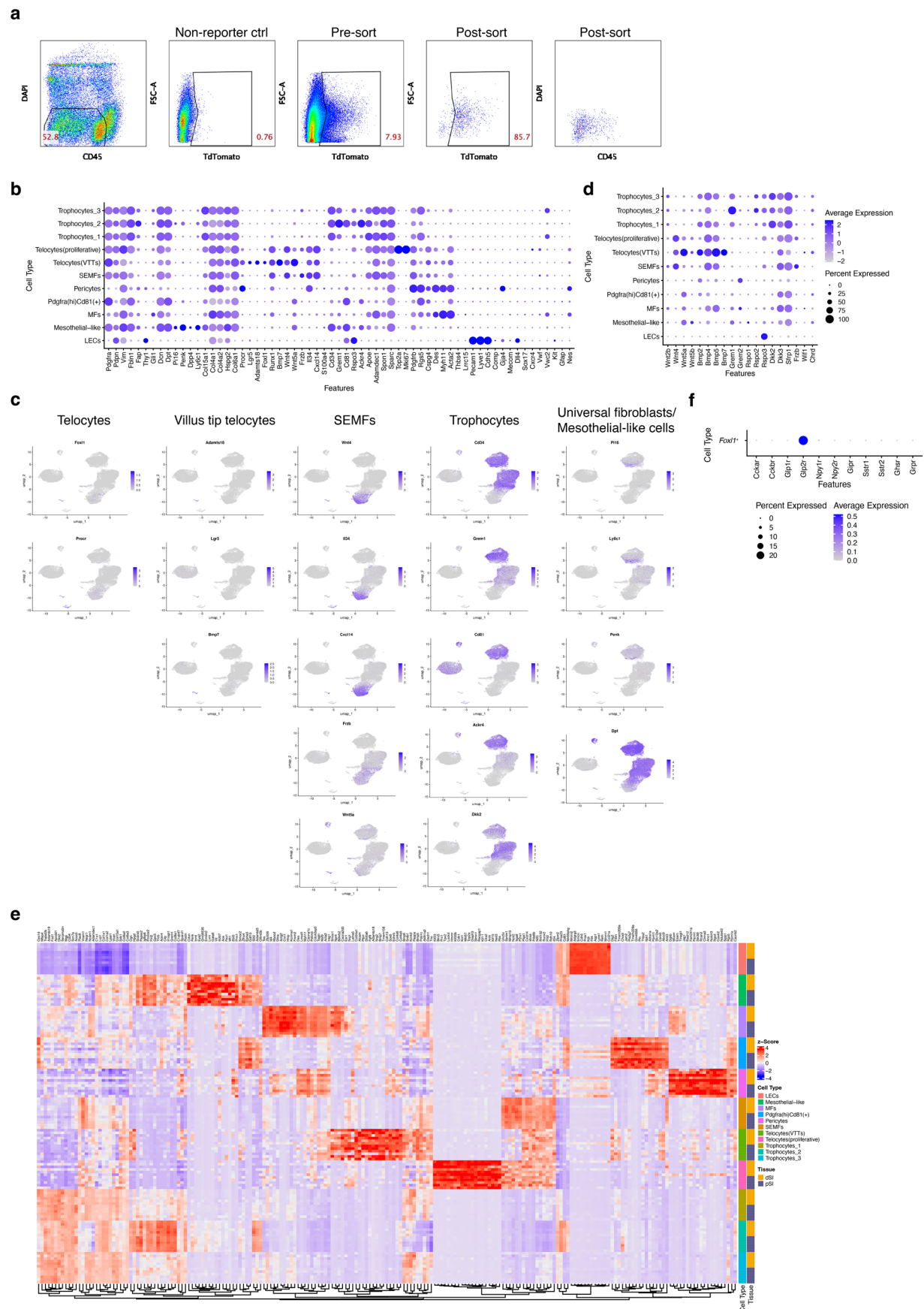
a, Representative imaging of CD201 and CD34 expression in proximal and distal small intestine of Flare-TSLP; *Lgr5*-eGFP dual reporter mice. Note that epithelial tuft cells can express *Lgr5*-eGFP. Scale bar, 100 μ m. **b**, Gating strategy for FACS sorting of CD45⁺CD31⁻ TSLP-tdTomato⁺CD201⁺, and CD45⁺CD31⁻ TSLP-tdTomato⁺CD34⁺ cells from small intestinal LP (siLP, whole tissue) of Flare-TSLP

mice. **c**, RT-qPCR analysis for *Tslp*, *Foxl1*, *Wnt5a*, *Lgr5*, *Bmp7*, *Grem1*, *Rspo3*, *Wnt2b*, *Bmp4* expression in sorted CD45⁺CD31⁻ TSLP-tdTomato⁺CD201⁺, and CD45⁺CD31⁻ TSLP-tdTomato⁺CD34⁺, CD45⁺CD31⁻CD201⁺, and CD45⁺CD31⁻CD34⁺ stromal cells from siLP (whole tissue) of Flare-TSLP mice. Data are biological replicates, $n \geq 3$. Data are representative of at least two independent experiments. Error bars indicate samples mean \pm SEM.



Extended Data Fig. 3 | Feeding increases intestinal TSLP. a–e, Feeding increases intestinal TSLP. **a**, Percentage of TSLP-tdTomato⁺ cells among CD45⁺ cells in ileal lamina propria by flow cytometric analysis in fasted mice 2 hr after oral gavage of food (refed) or water as volumetric control (sham). Biological replicates N = 20 for control group and N = 17 for refeeding group. Statistical analysis was performed using unpaired t test, **P < 0.01. Error bars indicate samples mean \pm SEM. **b**, ELISA of TSLP protein recovered from proximal jejunal tissue homogenates after oral gavage of food at designated times in fasted wild type (WT) mice. Biological replicates N = 3 per time point. Error bars indicate samples mean \pm SEM. **c**, ELISA of TSLP protein recovered from distal ileal tissue homogenates from fasted WT

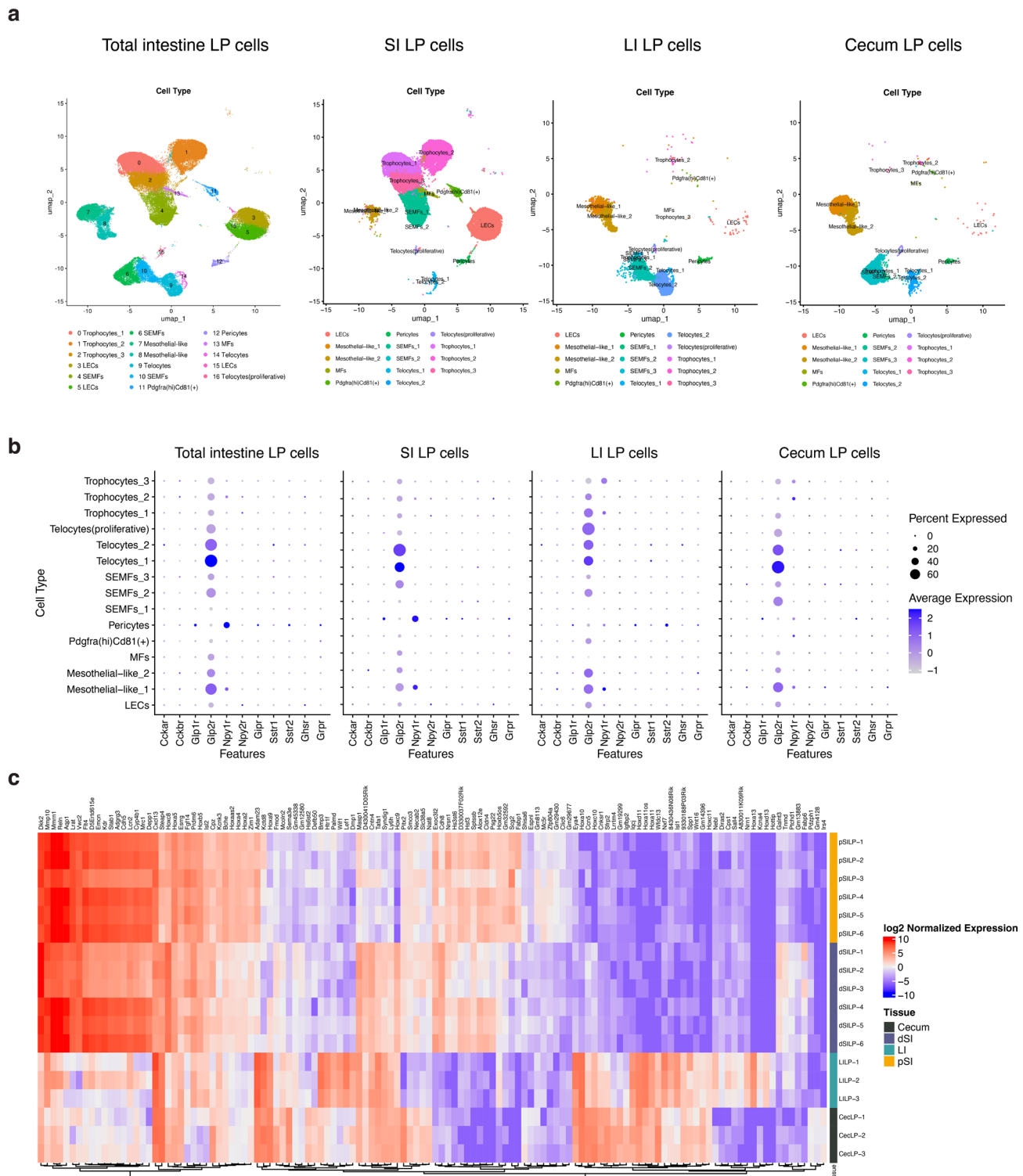
mice 2 hr after oral food gavage. Biological replicates N = 3–4 per time point. Statistical analysis was performed using ANOVA with correction for multiple comparisons. Error bars indicate samples mean \pm SEM. **d**, ELISA of TSLP protein recovered from proximal jejunal tissue homogenates after oral gavage 2 hr in Pou2f3^{-/-} or WT mice after overnight fasting. Biological replicates N = 3–7 each column group. Error bars indicate samples mean \pm SEM. **e**, ELISA of TSLP protein recovered from distal ileal tissue homogenates after oral gavage 2 hr in Pou2f3^{-/-} or WT mice after overnight fasting. Biological replicates N = 3–4 each column group. Error bars indicate samples mean \pm SEM.



Extended Data Fig. 4 | See next page for caption.

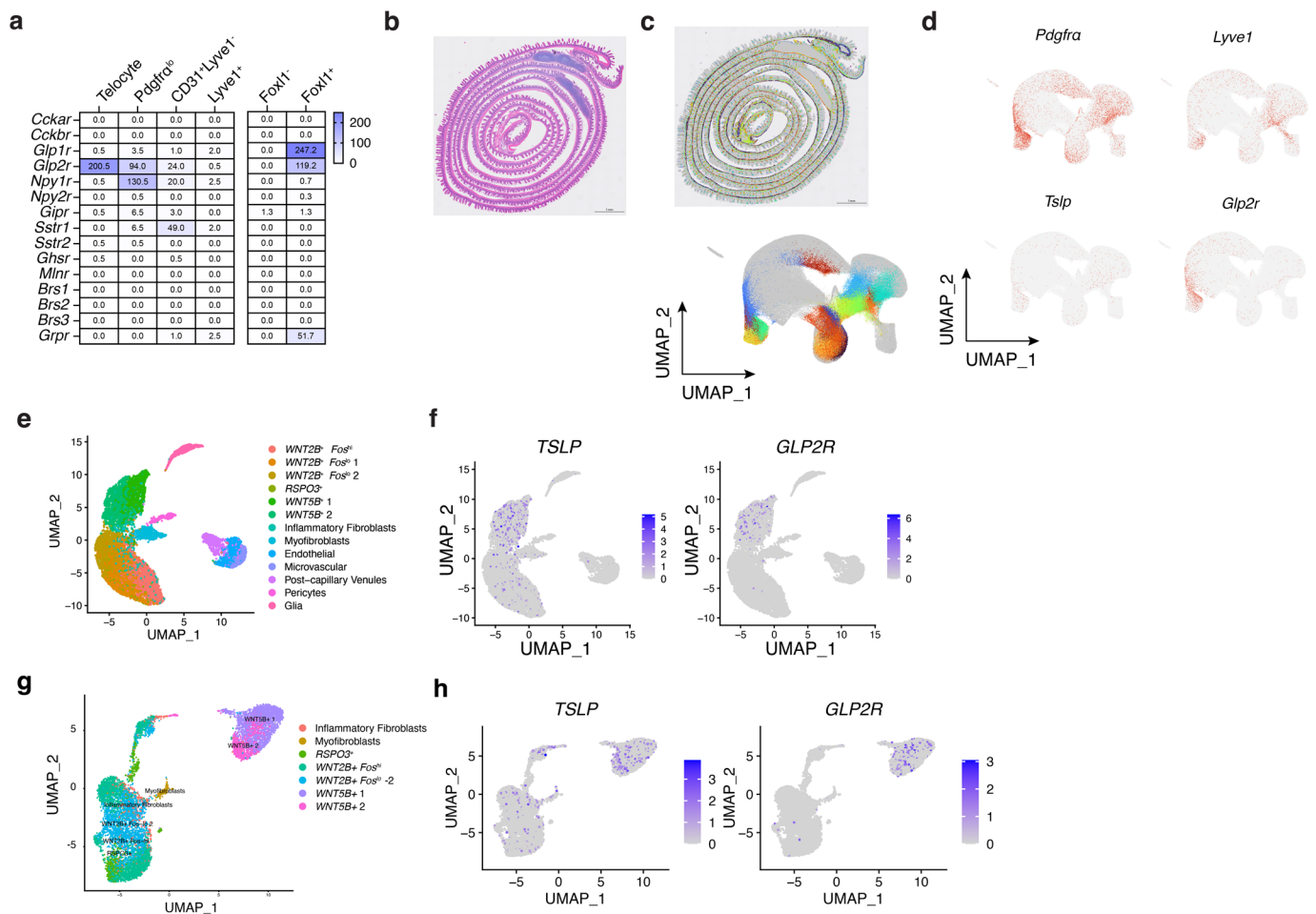
Extended Data Fig. 4 | Single cell RNA sequencing (scRNA-seq) analyses of TSLP-Tdtomato⁺ cells from mouse small intestinal lamina propria (siLP). **a**, Gating strategy for FACS sorting of TSLP⁺ tdTomato⁺ cells from intestinal LP of Flare-TSLP mice. **b–f**, scRNA-seq analysis of pooled proximal and distal siLP TSLP⁺ tdTomato⁺ cells. **b**, Expression levels of marker genes for annotated cell types. **c**, Feature plots represent marker genes for telocytes, VTTs, SEMFs, trophocytes and universal fibroblasts/mesothelial-like cells. **d**, Expression

levels of WNT and BMP signaling molecules for annotated cell types. **e**, Heatmap representation of top 20 most significantly changed genes including both directions from differential gene expression (DE) analysis between cell types in the proximal and distal small intestine. Columns were manually ordered by cell type. Plot shows z-scores. **f**, Expression levels of gut hormone receptors in *Foxl1*⁺ telocytes.



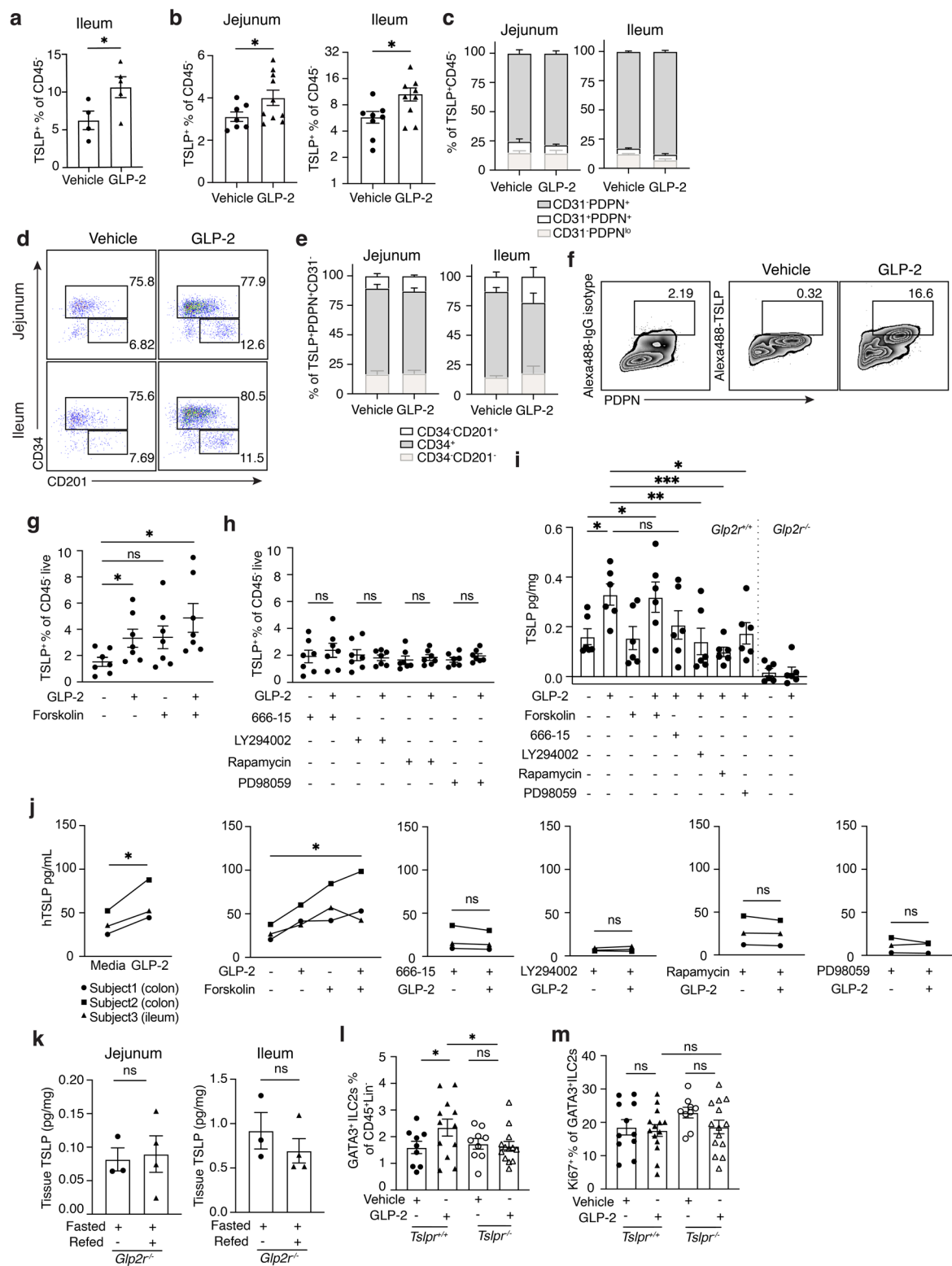
Extended Data Fig. 5 | scRNA-seq analyses of TSLP-TdTomato⁺ cells from mouse intestinal lamina propria (continued). **a**, UMAPs representing cell clustering in pooled cells (total intestine LP cells), siLP, LILP and cecum LP samples. Unsupervised cell clustering analysis revealed 17 clusters based on differential gene expression. Annotation was conducted according to dominant cluster-specific genes, while also considering existing stromal subsets. Subsets of fibroblasts were categorized as SEMFs (telocytes_2, SEMFs_1, SEMFs_2), trophocytes (trophocytes_1, trophocytes_2, trophocytes_3), and recently identified *Pdgfra*^{hi} universal fibroblasts or mesothelial-like cells (mesothelial-like_1, mesothelial-like_2). We also identified myofibroblasts (MFs), and an undefined cluster characterized by *Pdgfra*^{hi} *Cd81*⁺ *Ptn*⁺ fibroblasts.

Additional cells included LECs and pericytes. Some cell types were enriched in specific tissues; for instance, trophocytes, *Pdgfra*^{hi} *Cd81*⁺ cells, and LECs were primarily found in small intestine samples, while SEMFs_2 were more abundant in the large intestine and cecum but not in small intestine. Telocytes_2 were mostly enriched in the large intestine with few minimal in cecum and none in small intestine. **b**, Gut hormone receptor expression in total pooled intestine LP cells, LILP and cecum LP samples. **c**, Heatmap representing top 20 fold-change genes including both directions between tissue comparisons in DE analysis. Plot shows log₂ normalized expression. The transcriptomic signatures of TSLP-TdTomato⁺ cells were similar between proximal and distal small intestine, as well as between large intestine and cecum.



Extended Data Fig. 6 | scRNA-seq analyses of publicly available mouse and human intestinal stromal cell datasets. a, Heatmaps representing expression levels of gut hormone receptors in PDGFRa^{hi} telocytes¹⁵, PDGFRa^{lo} trophocytes¹⁵, CD31⁺LYVE1⁺ endothelial cells¹⁵, LYVE1⁺ LECs¹⁵, FoxI1⁺ telocytes¹⁶ and FoxI1⁺ stromal cells¹⁶ from mouse datasets. **b-d**, 10x Genomics published Visium mouse small intestine dataset⁵⁹. **b**, Hematoxylin and eosin staining of the intestinal tissue specimen. **c**, Color-coded cell clusters in the lamina propria (LP) shown

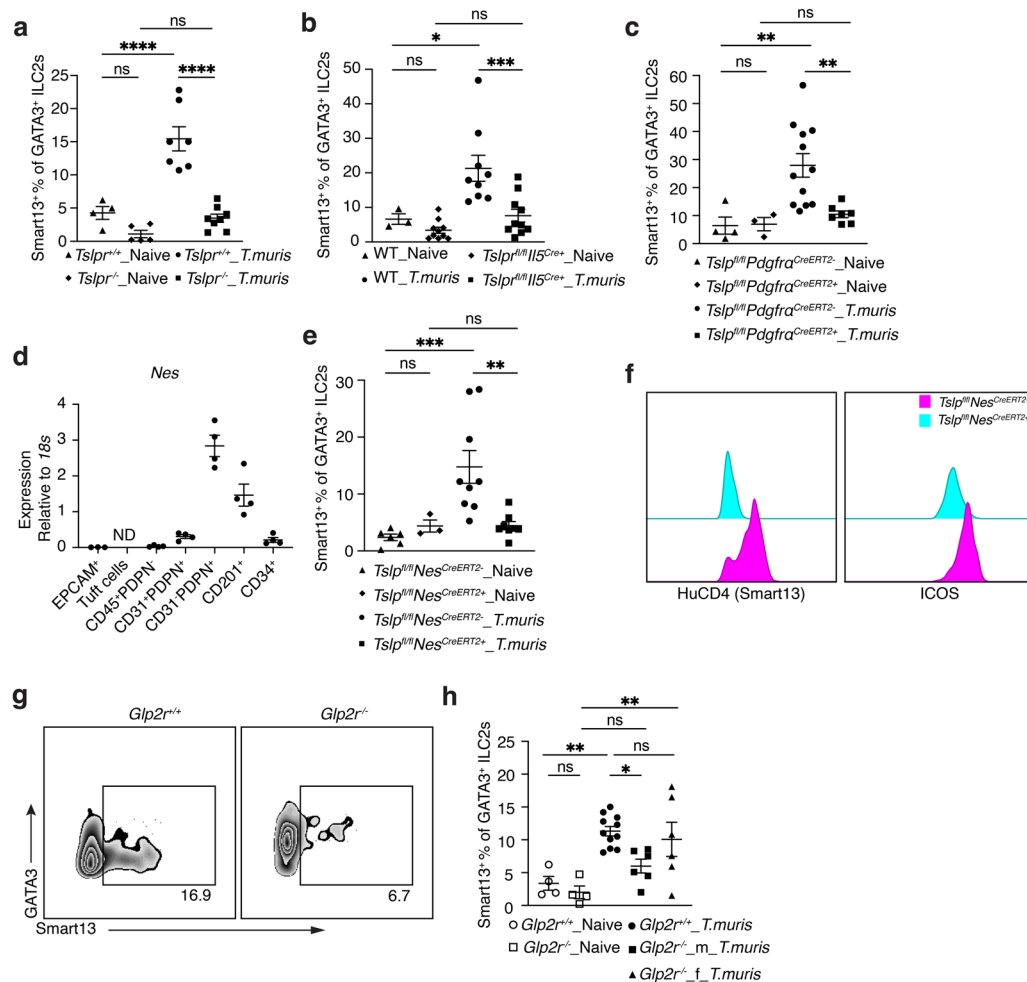
in situ (top) and on a UMAP (bottom). **d**, UMAP feature plots representing gene expression level and distribution of *Pdgfra*, *Lyve1*, *Tslp* and *Glp2r*. *Tslp* and *Glp2r* expression enriched in the stromal populations of intestinal LP. **e-h**, Analyses of human intestinal datasets^{24,25}. **e**, UMAP representing cell clustering of human dataset²⁴. **f**, Feature plots of *TSLP* (left) and *GLP2R* (right) expression levels. **g**, UMAP representing cell clustering of human dataset²⁵. **h**, Feature plots of *TSLP* (left) and *GLP2R* (right) expression levels.



Extended Data Fig. 7 | See next page for caption.

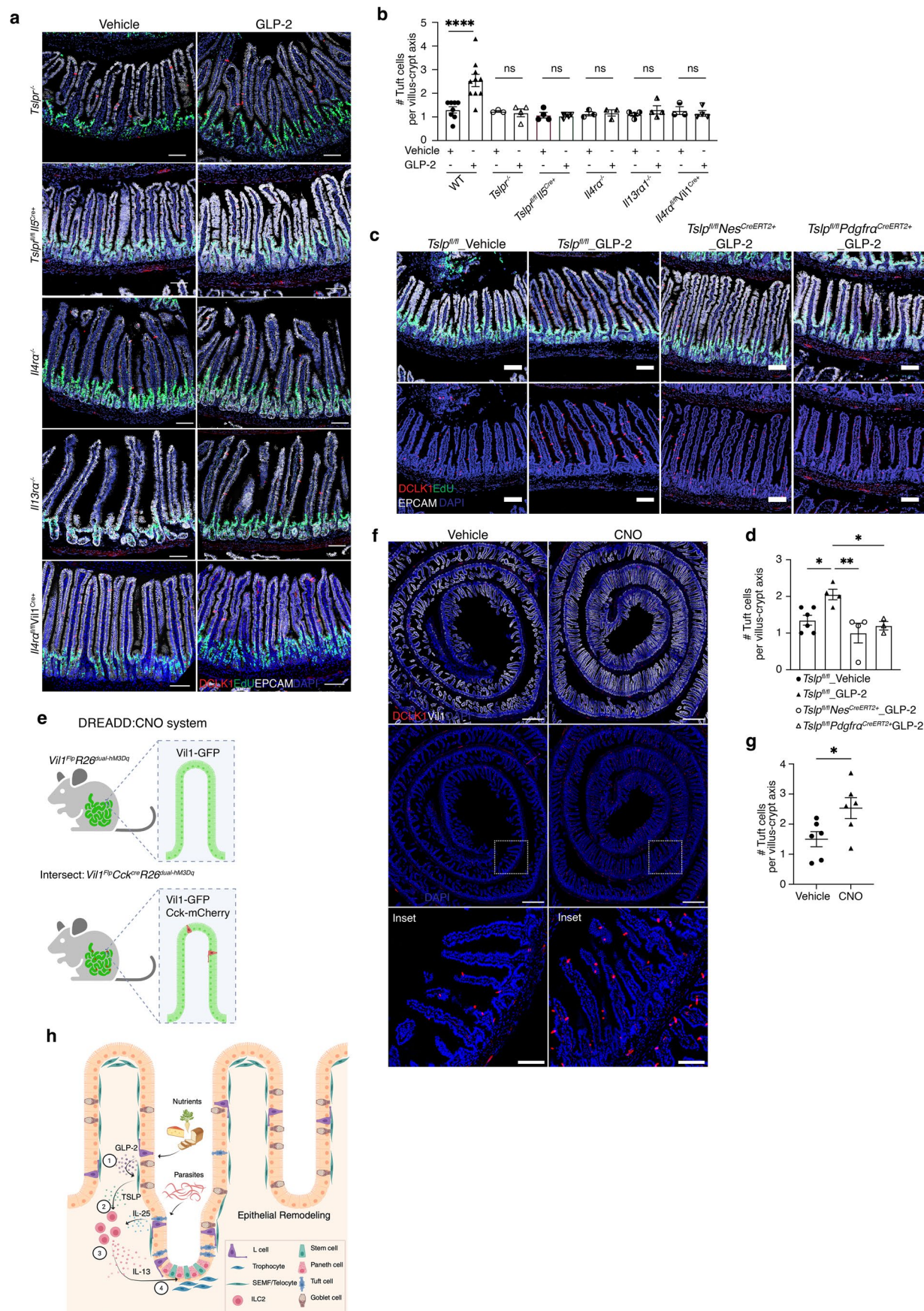
Extended Data Fig. 7 | L cell GLP-2 drives GLP-2R- and TSLP-dependent ILC2 activation. **a**, Flow cytometric analysis of TSLP-tdTomato⁺ cells in ileal lamina propria (LP) after 3 daily GLP-2[Gly2] injections. Biological replicates N = 4-5 for each group. **b-e** Flow cytometric analysis of TSLP-tdTomato⁺ cells in jejunal and ileal lamina propria (LP) at 2 hr after GLP-2[Gly2] administration. **b**, Percentage of total TSLP-tdTomato⁺ cells among CD45⁺ cells in jejunal (**left**) and ileal LP (**right**). Biological replicates N = 7-10 (left) and N = 8-9 (right) for each group. **c**, Proportion of CD31⁺PDPN⁺ LECs, CD31⁺PDPN⁺ and CD31⁺PDPN^{lo} stromal populations within TSLP-tdTomato⁺ cells in jejunum (**left**) and in ileum (**right**). Biological replicates N = 3-4 each group. **d-e**, Characterization of CD201⁺ and CD34⁺ cells within CD45⁺ PDPN⁺CD31⁺ TSLP-tdTomato⁺ stromal cells. **d**, Representative flow cytometric analysis. **e**, Quantification. Biological replicates N = 3-4 each column group. **f**, Intracellular antibody staining for TSLP protein in sorted CD45⁺CD31⁺TSLP-tdTomato⁺ stromal cells from small intestinal LP 4 hr after incubation with 10 µg/ml GLP-2[Gly2] with Brefeldin A *in vitro*. **g-h**, Mouse fibroblasts treated with agonists (**g**) and/or inhibitors (**h**). TSLP levels were assessed by flow cytometric analysis using frequency of TSLP⁺ cells among total

cultured fibroblasts characterized by CD45⁺PDPN⁺. Biological replicates N = 7 each group. **i**, Mouse intestinal explants treated with agonists and/or inhibitors. TSLP protein levels in the supernatants were measured by ELISA. Biological replicates total N = 77. **j**, TSLP protein levels in supernatants of human fibroblast cultures treated with agonists and/or inhibitors. Paired t-tests were performed for statistical analysis. Biological replicates N = 3. **k**, TSLP protein recovered from jejunal and ileal tissue homogenates from fasted *Glp2r*^{-/-} mice 2 hr after food gavage. Biological replicates N = 3-4 each group. **l-m**, Flow cytometric analysis of ILC2s in jejunal LP in *Tslpr*^{+/+} and *Tslpr*^{-/-} mice after 3 daily injections of GLP-2[Gly2]. **l**, Percentage of ILC2s among Lin⁻CD45⁺ cells. **m**, Quantification of percentage of Ki67⁺ cells among ILC2s. ILC2s were gated on CD45⁺Lin⁻GATA3⁺ cells. Biological replicates N = 9-14 for each group. Graphs represent data pooled from at least two experiments. Error bars indicate samples mean ± SEM. Unless specified, unpaired t-test was used for analyses of difference between 2 groups. ANOVA test was used for analyzing experiments with >2 groups adjusted for multiple comparisons. *P < 0.05, **P < 0.01, ***P < 0.001.



Extended Data Fig. 8 | *T. muris* infection drives stromal TSLP-dependent ILC2 activation. **a-c**, Flow cytometric analysis of ILC2s in large intestinal lamina propria (LILP) on day 21 post *T. muris* infection. Percentage of IL-13 (Smart13)⁺ ILC2 among ILC2s in LILP in *Tslpr*^{-/-} or *Tslpr*^{+/+} mice, biological replicates N = 4-8 each column group, ****P < 0.0001 (a), *Tslpr*^{R/R} *Il5*^{Cre/+} or WT controls, biological replicates N = 3-10 each column group, *P < 0.05, ***P < 0.001 (b), and *Tslpr*^{R/R} *Pdgfra*^{CreERT2/+} or *Tslpr*^{R/R} controls, biological replicates N = 3-12 each column group, **P < 0.01 (c). ILC2s were gated on Lin⁻CD45⁺GATA3⁺ cells. **d**, RT-qPCR analysis for *Nes* expression in sorted cells from the small intestine, including CD45⁺EPCAM⁺ epithelial cells, tuft cells, CD45⁺PDPN⁺ hematopoietic cells, CD45⁺CD31⁺PDPN⁺ LECs, CD45⁺CD31⁺PDPN⁺ CD201⁺, and CD45⁺CD31⁺CD34⁺ stromal cells. Biological replicates N = 3-4 each column group. **e-f**, Flow cytometric analyses of LILP ILC2s in *Tslpr*^{R/R} *Nes*^{CreERT2/+} or control mice on

21 d.p.i. Percentage of IL-13 (Smart13)⁺ ILC2 among ILC2s in LILP in *Tslpr*^{R/R} *Nes*^{CreERT2/+} or control (e) and the expression levels of ILC2s activation markers including HuCD4 (Smart13) and ICOS were lower in *Tslpr*^{R/R} *Nes*^{CreERT2/+} mice than controls (f). ILC2s were gated on Lin⁻CD45⁺GATA3⁺ cells. Biological replicates N = 3-9 each column group, **P < 0.01, ***P < 0.001. **g-h**, Flow cytometric analysis of LILP ILC2s in *Glp2r*^{-/-} and *Glp2r*^{+/+} mice 21 d.p.i. Flow charts represent IL-13 (Smart13) levels (g) and percentage of IL-13 (Smart13)⁺ ILC2s among total ILC2s in LILP (h). ILC2s were gated on Lin⁻CD45⁺GATA3⁺ cells. Biological replicates N = 4-11 each column group, *P < 0.05, **P < 0.01. Data are representative of at least two independent experiments. Statistical analyses were performed using ANOVA with correction for multiple comparisons. Error bars indicate samples mean ± SEM.



Extended Data Fig. 9 | See next page for caption.

Extended Data Fig. 9 | GLP-2 drives ILC2-dependent amplification of the tuft cell circuit. **a**, Representative imaging of jejunal tuft cells in *Tslpr*^{-/-}, *Tslpr*^{R/R}, *Il5*^{Cre+}, *Il4ra*^{-/-}, *Il13ra1*^{-/-} and *Il4ra*^{R/R}*Vil1*^{Cre+} mice. Green, EdU; Red, DCLK1; White, EPCAM; Blue, DAPI. 20x objective. Scale bar, 100 μm. **b**, Quantification of jejunal tuft cells after 3 daily GLP-2[Gly2] injections in WT, *Tslpr*^{-/-}, *Tslpr*^{R/R}*Il5*^{Cre+}, *Il4ra*^{-/-}, *Il13ra1*^{-/-} and *Il4ra*^{R/R}*Vil1*^{Cre+} mice. For tuft cell quantification, images were acquired using 20x objective and total DCLK1⁺EPCAM⁺ cells in each field were counted and normalized by total number of villus-crypt axes in the field. Each dot represents a mouse, N = 3-10 each column. Statistical analysis was performed using ANOVA with correction for multiple comparisons, ****P < 0.0001. Error bars indicate samples mean ± SEM. **c**, Representative imaging of jejunal tuft cells in *Tslp*^{R/R}, *Tslp*^{R/R}*Nes*^{CreERT2+}, and *Tslp*^{R/R}*Pdgfra*^{CreERT2+} mice. Green, EdU; Red, DCLK1; White, EPCAM; Blue, DAPI. 20x objective. Scale bar, 100 μm. **d**, Quantification of jejunal tuft cells after 3 daily GLP-2[Gly2] injections in *Tslp*^{R/R}, *Tslp*^{R/R}*Nes*^{CreERT2+}, and *Tslp*^{R/R}*Pdgfra*^{CreERT2+} mice. Each dot represents a mouse, N = 3-6. Statistical analysis was performed using ANOVA with correction for multiple comparisons, *P < 0.05,

P < 0.01. Error bars indicate samples mean ± SEM. **e, Schematic illustrating the *Vil1*^{Flp}*Cck*^{Cre}*R26*^{Dual-hM3Dq} DREADD and CNO system. **f-g**, Quantification of jejunal tuft cells in *Vil1*^{Flp}*Cck*^{Cre}*R26*^{Dual-hM3Dq} mice after administration of clozapine N-oxide (CNO) or vehicle control. Each dot represents a mouse. **f**, Representative imaging of tuft cells. Red, DCLK1; White, *Vil1*-GFP; Blue, DAPI. 20x objective. Scale bar, 100 μm. **g**, Quantification of tuft cells. Two experiment repeats, each dot represents a mouse, N = 6 for each column group. Statistical analysis was performed using two-tailed unpaired t-test, *P < 0.05. Error bars indicate samples mean ± SEM. **h**, Proposed model for the mechanism of GLP-2 – TSLP – ILC2 signaling pathway in promoting intestine tissue remodeling. GLP-2 from L cells (1) stimulates telocytes to produce TSLP (2) that activates ILC2s (3), thus linking nutrient ingestion and IL-13-mediated amplification of epithelial tuft cells. (4) *N. brasiliensis* short circuits the physiologic pathway by directly stimulating tuft cells to release IL-25 that activates ILC2s and drives the ILC2–tuft cell circuit amplification. Panels **e** and **h** created using BioRender.com.

Reporting Summary

Nature Portfolio wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Portfolio policies, see our [Editorial Policies](#) and the [Editorial Policy Checklist](#).

Please do not complete any field with "not applicable" or n/a. Refer to the help text for what text to use if an item is not relevant to your study.

For final submission: please carefully check your responses for accuracy; you will not be able to make changes later.

Statistics

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

n/a Confirmed

- ☐ ☒ The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
- ☐ ☒ A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
- ☐ ☒ The statistical test(s) used AND whether they are one- or two-sided
Only common tests should be described solely by name; describe more complex techniques in the Methods section.
- ☒ ☐ A description of all covariates tested
- ☐ ☒ A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
- ☐ ☒ A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
- ☐ ☒ For null hypothesis testing, the test statistic (e.g. F , t , r) with confidence intervals, effect sizes, degrees of freedom and P value noted
Give P values as exact values whenever suitable.
- ☒ ☐ For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
- ☒ ☐ For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
- ☒ ☐ Estimates of effect sizes (e.g. Cohen's d , Pearson's r), indicating how they were calculated

Our web collection on [statistics for biologists](#) contains articles on many of the points above.

Software and code

Policy information about [availability of computer code](#)

Data collection

Single cell RNA sequencing (sc-RNAseq) and cell hashing

Live tdTomato+ cells were purified using FACS and labeled with unique barcoded H-2/CD45 TotalSeq™-B anti-mouse hashtag antibodies (Biolegend Cat. 155831, 155833, 155835, 155837, 155839, 155841) for multiplexing. The cells were processed according to the manufacturer's instructions using the Chromium™ Next GEM Single Cell Kit v3.1 (10X Genomics, Cat. 1000269). Briefly, cell counts were performed using a hemacytometer and the desired number of cells was injected into microfluidic chips (10X Genomics, Cat. 1000127) to create Gel Beads-in-Emulsion (GEMs) with the 10X Chromium controller. Reverse transcription (RT) was performed on the GEMs followed by purification and amplification of the RT products. Cell multiplexing oligo (CMO) DNA was separated from cDNA using size selection with SPRI select beads (Beckman Coulter, Cat. B23318). Gene expression (GEX) libraries and CMO libraries were generated separately and profiled using the Bioanalyzer High Sensitivity DNA kit (Agilent Technologies, Cat. 5067-4626). Prior to sequencing, the GEX and corresponding CMO libraries were mixed in a 4:1 ratio. Sequencing of the mixed libraries was conducted on the Illumina NovaSeq X platform.

Data analysis

sc-RNAseq data analysis

Sample demultiplexing, alignments and UMI counts were performed using the 10x Genomics Cell Ranger pipeline (v 8.1.0) based on the latest mouse genome (mm39) with a custom reference to include the tdTomato sequence. Doublets were removed using the DoubletFinder (v2.0.4) algorithm and ambient RNA was eliminated with the SoupX algorithm (v1.6.2) using default parameters. After preprocessing datasets were analyzed using the Seurat single-cell analysis pipeline (v5.0.3) in R (v4.3.1). Quality control involved excluding low-quality cells based on the following criteria: nFeature_RNA between 1000 and 7500, nCount_RNA between 2500 and 40000, and mitochondrial contamination below 10%. Following filtering data were normalized using the NormalizeData function with the "LogNormalize" method and the 2000 most variable genes were identified and scaled before performing PCA. Thirty principal components were used for graph-based clustering (resolution = 0.5) followed by UMAP dimensionality reduction.

An additional filtering step based on tdTomato expression removed contaminant clusters with fewer than 0.01% tdTomato+ cells and fewer than 5 absolute tdTomato+ cells. Initial cell type annotation was conducted automatically using scType (58) followed by manual refinement. Differential gene expression (DE) analysis between cell types in the small intestine samples were conducted using Wilcoxon rank sum tests as implemented in the Seurat function FindAllMarkers. For DE analysis between tissues, counts from each sample were summed and followed by pseudobulk DE analysis using limma-voom.

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio [guidelines for submitting code & software](#) for further information.

Data

Policy information about [availability of data](#)

All manuscripts must include a [data availability statement](#). This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A description of any restrictions on data availability
- For clinical datasets or third party data, please ensure that the statement adheres to our [policy](#)

All raw and processed scRNA-seq data generated in this study have been deposited in the NCBI Gene Expression Omnibus (GEO) under accession numbers GSE280635. All the other data supporting the findings of this study are available.

Research involving human participants, their data, or biological material

Policy information about studies with [human participants or human data](#). See also policy information about [sex, gender \(identity/presentation\), and sexual orientation](#) and [race, ethnicity and racism](#).

Reporting on sex and gender	Primary cell lines established from gut tissues of two male and one female human subjects.
Reporting on race, ethnicity, or other socially relevant groupings	N/A
Population characteristics	N/A
Recruitment	Patients underwent colonoscopy or sigmoidoscopy for non-inflammatory indications (e.g., colorectal cancer screening) and were referred as healthy controls (HC). Baseline demographic information for the study participants are provided in Supplementary Table S1. Patients consented to publish de-identified patient demographics including age at the time of sample collection, sex, diagnosis, and medical center. Demographic options were defined by the investigators and participants chose their classifications.
Ethics oversight	The study was conducted in accordance with the principles of the Declaration of Helsinki and was approved by the Institutional Review Board of the University of California, San Francisco (19-27302). All participants provided written informed consent before inclusion.

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

Field-specific reporting

Please select the one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.

☒ Life sciences ☐ Behavioural & social sciences ☐ Ecological, evolutionary & environmental sciences

For a reference copy of the document with all sections, see nature.com/documents/nr-reporting-summary-flat.pdf

Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

Sample size	Describe how sample size was determined, detailing any statistical methods used to predetermine sample size OR if no sample-size calculation was performed, describe how sample sizes were chosen and provide a rationale for why these sample sizes are sufficient.
Data exclusions	Describe any data exclusions. If no data were excluded from the analyses, state so OR if data were excluded, describe the exclusions and the rationale behind them, indicating whether exclusion criteria were pre-established.
Replication	Describe the measures taken to verify the reproducibility of the experimental findings. If all attempts at replication were successful, confirm this OR if there are any findings that were not replicated or cannot be reproduced, note this and describe why.
Randomization	Describe how samples/organisms/participants were allocated into experimental groups. If allocation was not random, describe how covariates were controlled OR if this is not relevant to your study, explain why.

Describe whether the investigators were blinded to group allocation during data collection and/or analysis. If blinding was not possible, describe why OR explain why blinding was not relevant to your study.

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Materials & experimental systems

- | | |
|-------------------------------------|---|
| n/a | Involvement in the study |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> Antibodies |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> Eukaryotic cell lines |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Palaeontology and archaeology |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> Animals and other organisms |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Clinical data |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Dual use research of concern |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Plants |

Methods

- | | |
|--------------------------|--|
| n/a | Involvement in the study |
| <input type="checkbox"/> | <input type="checkbox"/> ChIP-seq |
| <input type="checkbox"/> | <input checked="" type="checkbox"/> Flow cytometry |
| <input type="checkbox"/> | <input type="checkbox"/> MRI-based neuroimaging |

Antibodies

Antibodies used

All antibodies used have been detailed described in the manuscript.

Validation

Antibodies were purchased from reliable commercial sources.

Antibodies	Suppliers	Catalogs	Dilution	Website
Brilliant Violet 421™ anti-mouse CD19 Antibody	BioLegend	115549	1:200	https://www.biolegend.com/fr-lu/products/brilliant-violet-421-anti-mouse-cd19-antibody-7160
Brilliant Violet 421™ anti-mouse TER-119/Erythroid Cells Antibody	BioLegend	116234	1:200	https://www.biolegend.com/fr-lu/products/brilliant-violet-421-anti-mouse-ter-119-erythroid-cells-antibody-7259
Brilliant Violet 421™ anti-mouse Ly-6G/Ly-6C (Gr-1) Antibody	BioLegend	108445	1:200	https://www.biolegend.com/fr-lu/products/brilliant-violet-421-anti-mouse-ly-6g-ly-6c-gr-1-antibody-7201
Brilliant Violet 605™ anti-human/mouse/rat CD278 (ICOS) Antibody	BioLegend	313538	1:100	https://www.biolegend.com/fr-lu/products/brilliant-violet-605-anti-human-mouse-rat-cd278-icos-antibody-14371
Brilliant Violet 421™ anti-mouse/human CD11b Antibody	BioLegend	101251	1:200	https://www.biolegend.com/fr-lu/products/brilliant-violet-421-anti-mouse-human-cd11b-antibody-7163
Pacific Blue™ anti-mouse CD11c Antibody	BioLegend	117322	1:200	https://www.biolegend.com/fr-lu/products/pacific-blue-anti-mouse-cd11c-antibody-3864
Pacific Blue™ anti-mouse CD49b (pan-NK cells) Antibody	BioLegend	108918	1:200	https://www.biolegend.com/fr-lu/products/pacific-blue-anti-mouse-cd49b-pan-nk-cells-antibody-6571
Brilliant Violet 421™ anti-mouse CD335 (NKP46) Antibody	BioLegend	137612	1:200	https://www.biolegend.com/fr-lu/products/brilliant-violet-421-anti-mouse-cd335-nkp46-antibody-7506
Brilliant Violet 421™ anti-mouse NK-1.1 Antibody	BioLegend	108741	1:200	https://www.biolegend.com/fr-lu/products/brilliant-violet-421-anti-mouse-nk-1-1-antibody-7150
Brilliant/Violet 421™ anti-mouse TCR γ/δ Antibody	BioLegend	118120	1:200	https://www.biolegend.com/fr-lu/products/brilliant-violet-421-anti-mouse-tcr-gamma-delta-antibody-7249
Pacific Blue™ anti-mouse FcεRIα Antibody	BioLegend	134314	1:200	https://www.biolegend.com/fr-lu/products/pacific-blue-anti-mouse-fcepsilonrialpha-antibody-6703
Pacific Blue™ anti-mouse F4/80 Antibody	BioLegend	123124	1:200	https://www.biolegend.com/fr-lu/products/pacific-blue-anti-mouse-f4-80-antibody-4075
BUV395 Rat Anti-Mouse CD45	BD	564279	1:200	https://www.bdbiosciences.com/en-us/products/reagents/flow-cytometry-reagents/research-reagents/single-color-antibodies-ruo/buv395-rat-anti-mouse-cd45.564279?tab=product_details
PE/Cyanine7 anti-mouse/human KLRG1 (MAFA) Antibody	BioLegend	138416	1:100	https://www.biolegend.com/fr-lu/products/pe-cyanine7-anti-mouse-human-klrg1-mafa-antibody-8312
APC anti-human CD4 Antibody	BioLegend	300514	1:50	https://www.biolegend.com/fr-lu/products/apc-anti-human-cd4-antibody-823
Ki-67 Monoclonal Antibody (SolA15), FITC, eBioscience™	Invitrogen	11-5698-82	1:1000	https://www.thermofisher.com/antibody/product/Ki-67-Antibody-clone-SolA15-Monoclonal/11-5698-82
APC anti-mouse CD201 (EPCR) Antibody	BioLegend	141506	1:200	https://www.biolegend.com/fr-lu/products/apc-anti-mouse-cd201-ePCR-antibody-7886
PerCP/Cyanine5.5 anti-mouse CD31 Antibody	BioLegend	102420	1:200	https://www.biolegend.com/fr-lu/products/percp-cyanine5-5-anti-mouse-cd31-antibody-6668
Brilliant Violet 421™ anti-mouse CD31 Antibody	BioLegend	102424	1:200	https://www.biolegend.com/fr-lu/products/brilliant-violet-421-anti-mouse-cd31-antibody-8599
Brilliant Violet 605™ anti-mouse CD140α Antibody	BioLegend	135916	1:100	https://www.biolegend.com/fr-lu/products/brilliant-violet-605-anti-mouse-cd140a-antibody-15109

Alexa Fluor® 488 anti-mouse LYVE1 Antibody	eBioscience s	53-0443-82	1:100	https://www.thermofisher.com/antibody/product/LYVE1-Antibody-clone-ALY7-Monoclonal/53-0443-82
Alexa Fluor® 488 anti-mouse Podoplanin Antibody	BioLegend	127406	1:100	https://www.biolegend.com/fr-lu/products/alexa-fluor-488-anti-mouse-podoplanin-antibody-4751
BV786 Rat anti-mouse CD34	BD	742971	1:100	https://www.bdbiosciences.com/en-us/products/reagents/flow-cytometry-reagents/research-reagents/single-color-antibodies-ruo/bv786-rat-anti-mouse-cd34.742971?tab=product_details
Alexa Fluor® 647 anti-mouse CD326 (Ep-CAM) Antibody	BioLegend	118212	1:100	https://www.biolegend.com/fr-lu/products/alexa-fluor-647-anti-mouse-cd326-ep-cam-antibody-4973
Alexa Fluor® 488 anti-mouse CD326 (Ep-CAM) Antibody	BioLegend	118210	1:100	https://www.biolegend.com/fr-lu/products/alexa-fluor-488-anti-mouse-cd326-ep-cam-antibody-4972
Alexa Fluor® 488 Rat IgG2a, κ Isotype Ctrl Antibody	BioLegend	400525	1:100	https://www.biolegend.com/fr-lu/products/alexa-fluor-488-rat-igg2a-kappa-isotype-ctrl-2682
Anti-DCAMKL1 antibody	Abcam	ab31704	1:500	https://www.abcam.com/en-us/products/primary-antibodies/dcamkl1-antibody-ab31704?srsId=AfmBOooWK-kIGb0WmC-vZVqchamVQrtivBIWv_xQ8Y77KO2gv6dH_Bdi
Mouse Podoplanin Antibody	R&D	AF3244	1:250	https://www.rndsystems.com/products/mouse-podoplanin-antibody_af3244
Mouse PDGFR alpha Antibody	R&D	MAB1062	1:250	https://www.rndsystems.com/products/mouse-pdgf-ralpha-antibody-189208_mab1062
Mouse EPCR (CD201) Antibody	R&D	AF2749	1:200	https://www.rndsystems.com/products/mouse-epcr-antibody_af2749
CD34 Monoclonal Antibody (RAM34), eBioscience™	ThermoFisher	14-0341-82	1:500	https://www.thermofisher.com/antibody/product/CD34-Antibody-clone-RAM34-Monoclonal/14-0341-82
Anti-Green Fluorescent Protein Antibody	Aveslabs	GFP-1020	1:500	https://www.antibodiesinc.com/products/anti-green-fluorescent-protein-antibody-gfp?srsId=AfmBOooLZXsckBI81Y2anClkRxJt3LRieoe8OCCXdFwR9xoVGy0PBNKh
Living Colors® DsRedPolyclonal Antibody	TakaraBio	632496	1:250	https://www.takarabio.com/products/antibodies-and-elisa/fluorescent-protein-antibodies/red-fluorescent-protein-antibodies?srsId=AfmBOornk_0PBiq4yY73d8T288enog67I0VqwUNtZ4I5jSFFBzVmZx6O
mCherry Monoclonal Antibody (16D7)	ThermoFisher	M11217	1:500	https://www.thermofisher.com/antibody/product/mCherry-Antibody-clone-16D7-Monoclonal/M11217
TSLP Monoclonal Antibody	Invitrogen	MA5-23779	1:100	https://www.thermofisher.com/antibody/product/TSLP-Antibody-clone-152640-Monoclonal/MA5-23779

Eukaryotic cell lines

Policy information about [cell lines and Sex and Gender in Research](#)

Cell line source(s)	Primary intestinal fibroblasts derived from healthy donors, 2 males and 1 female; generated and provided by Michael G. Kattah
Authentication	The primary human fibroblast cell lines were originally derived by our coauthors and not commercially purchased. The methodology of deriving the cell lines have been detailed described in the Methods section. All cell lines have been grown below passage 9. The cell lines have been tested by microscopy to verify fibroblast identity and transcriptomic profiling confirming fibroblast marker gene signatures.
Mycoplasma contamination	Negative for Mycoplasma
Commonly misidentified lines (See ICLAC register)	N/A

Animals and other research organisms

Policy information about [studies involving animals](#); [ARRIVE guidelines](#) recommended for reporting animal research, and [Sex and Gender in Research](#)

Laboratory animals	Mouse
Wild animals	N/A
Reporting on sex	Both female and male mice were used; sex-bias (if any) was reported in the manuscript.
Field-collected samples	N/A
Ethics oversight	UCSF Institutional Animal Care and Use Committee (IACUC); UCSF Laboratory Animal Resource Center (LARC)

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

Plants

Seed stocks	Report on the source of all seed stocks or other plant material used. If applicable, state the seed stock centre and catalogue number. If plant specimens were collected from the field, describe the collection location, date and sampling procedures.
Novel plant genotypes	Describe the methods by which all novel plant genotypes were produced. This includes those generated by transgenic approaches, gene editing, chemical/radiation-based mutagenesis and hybridization. For transgenic lines, describe the transformation method, the number of independent lines analyzed and the generation upon which experiments were performed. For gene-edited lines, describe the editor used, the endogenous sequence targeted for editing, the targeting guide RNA sequence (if applicable) and how the editor was applied.
Authentication	Describe any authentication procedures for each seed stock used or novel genotype generated. Describe any experiments used to assess the effect of a mutation and, where applicable, how potential secondary effects (e.g. second site T-DNA insertions, mosaicism, off-target gene editing) were examined.

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 <i>May remain private before publication.</i>	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	Standard live cell surface antibody staining and intracellular staining
Instrument	BD Fortessa; BD Aria; Cytex Aurora
Software	FlowJo

Cell population abundance

Reported in the manuscript

Gating strategy

Included in the supplementary

☒ 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)

Specify: functional, structural, diffusion, perfusion.

Field strength

Specify in Tesla

Sequence & imaging parameters

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.

Area of acquisition

State whether a whole brain scan was used OR define the area of acquisition, describing how the region was determined.

Diffusion MRI

☐ Used☐ Not used

Preprocessing

Preprocessing software

Provide detail on software version and revision number and on specific parameters (model/functions, brain extraction, segmentation, smoothing kernel size, etc.).

Normalization

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.

Normalization template

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.

Noise and artifact removal

Describe your procedure(s) for artifact and structured noise removal, specifying motion parameters, tissue signals and physiological signals (heart rate, respiration).

Volume censoring

Define your software and/or method and criteria for volume censoring, and state the extent of such censoring.

Statistical modeling & inference

Model type and settings

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).

Effect(s) tested

Define precise effect in terms of the task or stimulus conditions instead of psychological concepts and indicate whether ANOVA or factorial designs were used.

Specify type of analysis: ☐ Whole brain ☐ ROI-based ☐ Both

Statistic type for inference

Specify voxel-wise or cluster-wise and report all relevant parameters for cluster-wise methods.

(See [Eklund et al. 2016](#))

Correction

Describe the type of correction and how it is obtained for multiple comparisons (e.g. FWE, FDR, permutation or Monte Carlo).

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

Report the measures of dependence used and the model details (e.g. Pearson correlation, partial correlation, mutual information).

Graph analysis

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.).

Multivariate modeling and predictive analysis

Specify independent variables, features extraction and dimension reduction, model, training and evaluation metrics.