Phospho-regulation of ATOH1 Is Required for Plasticity of Secretory Progenitors and Tissue Regeneration

Graphical Abstract

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In Brief
Tomic et al. report that multisite phosphorylation of ATOH1 regulates the contribution of secretory progenitors to stem cell self-renewal in the small intestine and colon. With damage, the enhanced role of Atoh1+ progenitors in mediating tissue repair is ablated in mice expressing phosphomutant ATOH1 and overall tissue regeneration is impaired.

Highlights
- Atoh1+ progenitors contribute to the stem cell pool in homeostasis and regeneration
- Multisite phosphorylation of ATOH1 regulates the plasticity of secretory progenitors
- Loss of phosphorylation of ATOH1 reduces clonogenic capacity of Atoh1+ cells
- Phosphomutant ATOH1 mice are more susceptible to chemical colitis
Phospho-regulation of ATOH1 Is Required for Plasticity of Secretory Progenitors and Tissue Regeneration

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SUMMARY

The intestinal epithelium is largely maintained by self-renewing stem cells but with apparently committed progenitors also contributing, particularly following tissue damage. However, the mechanism of, and requirement for, progenitor plasticity in mediating pathological response remain unknown. Here we show that phosphorylation of the transcription factor Atoh1 is required for both the contribution of secretory progenitors to the stem cell pool and for a robust regenerative response. As confirmed by lineage tracing, Atoh1+ cells (Atoh1WTCreERT2 mice) give rise to multilineage intestinal clones both in the steady state and after tissue damage. In a phosphomutant Atoh1(9S/T-A)CreERT2 line, preventing phosphorylation of ATOH1 protein acts to promote secretory differentiation and inhibit the contribution of progenitors to self-renewal. Following chemical colitis, Atoh1+ cells of Atoh1(9S/T-A)CreERT2 mice have reduced clonogenicity that affects overall regeneration. Progenitor plasticity maintains robust self-renewal in the intestinal epithelium, and the balance between stem and progenitor fate is directly coordinated by ATOH1 multisite phosphorylation.

INTRODUCTION

Within the intestinal epithelium, cell generation occurs from phenotypically heterogenous stem cells residing at the base of glandular crypts (Vermeulen and Snippert, 2014). There is broad consensus that this heterogeneity reflects the combined behavior of active and reserve stem cells. The former dominates in homeostatic self-renewal and the latter following tissue damage. In homeostasis, rapidly cycling stem cells express the R-spondin receptor Lgr5. Reserve stem cell function is less defined and has been ascribed variously to a subset of quiescent Lgr5+ cells (Barriga et al., 2017; Buczacki et al., 2013), progenitors committed to different intestinal lineages (van Es et al., 2012; Tetteh et al., 2016), and cells dependent on alternate pathways for stem cell maintenance (Takeda et al., 2011; Tian et al., 2011).

It has been demonstrated previously that cells of the secretory lineage possess reserve stem cell function in the small intestine (SI) epithelium in homeostasis and following tissue damage (van Es et al., 2012; Ishibashi et al., 2018; Yan et al., 2017; Yu et al., 2018). Subsequent to Delta-like expression (from Dll1 or Dll4), the basic helix-loop-helix (bHLH) transcription factor Atoh1 is upregulated, an event required for the creation of all secretory lineages within the epithelium (Yang et al., 2001). Atoh1+ progenitors exhibit self-renewal and give rise to multilineage clones with higher frequency in homeostasis (Ishibashi et al., 2018) compared with previously described secretory Dll1+ progenitors (van Es et al., 2012). This observation highlights a significant contribution of Atoh1+ cells to the stem cell pool in the SI and colon. However, the mechanisms regulating intestinal plasticity and the nature of the relationship linking it to self-renewal remain unknown.

ATOH1 can be phosphorylated on multiple sites by cyclin-dependent kinases. Here we demonstrate that maintenance of the plasticity of committed secretory precursors allowing return to the stem compartment is dependent on the multisite phosphorylation of ATOH1, prevention of which inhibits Atoh1-mediated self-renewal and results in compromised regeneration following damage. We conclude that reversibility of the commitment to differentiate is dependent on post-translational control of ATOH1 and is required to maintain a robust stem cell population.

RESULTS

Atoh1+ Cells Show Stem Cell Activity

Initially, to determine the extent to which Atoh1-expressing cells support stem cell maintenance in homeostasis, we generated a mouse (Atoh1WTCreERT2) with an inducible CreERT2 downstream of the Atoh1 coding sequence (Figure S1A). Acute lineage tracing demonstrated that tdTomato (tdTom) reporter
expression 24 hr following a single pulse of tamoxifen was restricted to secretory cells within the SI and colonic epithelium (Figures 1A–1D; Figures S1B–S1G). Mature Paneth and goblet cells were positive for the reporter whereas enteroendocrine cells (EECs) were not; the latter observation confirms that Atoh1 expression is not maintained in mature enteroendocrine cells (Bjerknes et al., 2012; Sommer and Mostoslavsky, 2014). However, by 4 days post-tamoxifen, enteroendocrine cells were also labeled (Figure 1E), indicating an origin from a secretory precursor that expressed Atoh1. Tuft cells were also not labeled with tdTom (Figure 1F). Individual Paneth cells remained labeled 4 weeks post-induction, reflecting their longevity (Figure S1H). Similar results were found in the colon, and long-lived secretory cells were also identified (Figure S1I). By 30 days post-induction, cohesive patches of reporter-positive cells that occupied all or a significant portion of entire crypts were present (Figures 1G and 1H) and continued to be observed after several months (Figure S1J). Immunostaining established the presence of goblet, Paneth, enteroendocrine, and absorptive cells were positive for the reporter whereas enteroendocrine cells (EECs) were not; the latter observation confirms that Atoh1 expression is not maintained in mature enteroendocrine cells (Bjerknes et al., 2012; Sommer and Mostoslavsky, 2014). However, by 4 days post-tamoxifen, enteroendocrine cells were also labeled (Figure 1E), indicating an origin from a secretory precursor that expressed Atoh1. Tuft cells were also not labeled with tdTom (Figure 1F). Individual Paneth cells remained labeled 4 weeks post-induction, reflecting their longevity (Figure S1H). Similar results were found in the colon, and long-lived secretory cells were also identified (Figure S1I). By 30 days post-induction, cohesive patches of reporter-positive cells that occupied all or a significant portion of entire crypts were present (Figures 1G and 1H) and continued to be observed after several months (Figure S1J). Immunostaining established the presence of goblet, Paneth, enteroendocrine, and absorptive

Figure 1. Lineage Tracing of Atoh1+ Cells in Homeostasis and after Injury

(A–D) The tdTom reporter is detected in Muc2+ goblet cells in the SI (A), colon (B), and Lyz+ Paneth cells (C) but not in ChgA+ enteroendocrine cells 24 hr post-tamoxifen (D). Muc2, Mucin 2; Lyz, Lysozyme; ChgA, Chromogranin A.

(E) ChgA+ cells labeled with tdTom on day 4 after induction.

(F) Dclk1+ tuft cells are not labeled with tdTom at 24 hr.

(G and H) Reporter-positive clone in the SI (G) and colon (H) 30 days following tamoxifen.

(I–L) tdTom+ clones at 30 days are composed of alkaline phosphatase (Alpi+) enterocytes (I), Paneth cells (J), goblet cells (K), and enteroendocrine cells (L).

(M, P, and S) Schematic of induction and injury protocol: irradiation (M), azoxymethane (AOM) (P), and dextran sodium sulfate (DSS) (S).

(N) Representative pictures of SI whole-mounts containing labeled crypts (arrowheads) 30 days post-induction.

(O) Quantification of tdTom+ crypts in the SI (n = 4 for 0 Gy, n = 6 for 6 Gy [day 1], n = 4 for 6 Gy [day 5]).

(Q and T) Representative images of colonic crypts on day 30 post-tamoxifen and AOM (Q) or DSS treatment (T). Note the large tdTom+ regenerative multicrypt patches (MCPs) associated with 2% DSS treatment (T).

(R) Quantification of reporter-positive crypts in the colon (n = 6 for untreated, n = 5 for AOM-treated).

(U) Quantification of tdTom+ MCPs in untreated and DSS-treated colons (n = 3 for both groups).

Welch’s t test was used in (O) (mean ± SEM, ****p < 0.0001) and Mann-Whitney test in (R) (mean ± SEM, **p = 0.0087). Scale bars, 50 µm (A–L) and 100 µm (N, Q, and T). See also Figure S1.
cells within reporter-positive epithelium, confirming their multilinage composition (Figures 1I–1L). These patterns are identical to those arising from individual marked intestinal stem cells (Vermeulen et al., 2013) and demonstrate a clonal origin from Atoh1+ precursors.

Atoh1(WT)CreERT2; Rosa26 TdTom mice were then crossed onto Lgr5Gfp reporter mice to investigate co-expression of Atoh1 and the intestinal stem cell marker Lgr5. The expression of Atoh1 and the tdTom reporter was identified in 1%–2% of Lgr5+ (GFP+) cells (Figures S1K–S1O), representing a likely intermediate state in the commitment process and candidate clonogetic population. Together, these results confirm that Atoh1 is appropriately expressed in mature Paneth and goblet cells but not enteroendocrine cells and that a proportion of Atoh1+ progenitors are acting as long-term multipotential stem cells (Bjerknes et al., 2012; Sommer and Mostoslavsky, 2014; Ishibashi et al., 2018).

**Atoh1+ Cells Contribute Directly to Epithelial Regeneration**

The extent of reversibility of Atoh1+ cell commitment was studied in the context of irradiation-induced tissue damage. Irradiation given 1 day after tamoxifen generated an increased number of tdTom+ crypts at 30 days in the SI compared with unirradiated controls (16-fold increase, 2.37% versus 0.15%). This effect was abrogated when irradiation was given 5 days after tamoxifen (Figures 1M–1O), suggesting that regenerative potential is a property of progenitors arising de novo from the stem cell compartment and not of more mature secretory cells. Similarly, after targeted deletion of the bulk of Lgr5 stem cells using a diphtheria toxin approach (Figures S1P and S1Q), there was a 30-fold increase in the number of clones observed (Figures S1R and S1S).

Adapting the assay to perform a similar analysis for the colonic epithelium and to circumvent that tissue’s known radio-resistance (Cai et al., 1997), mice were treated with the colon-specific carcinogen azoxymethane (AOM) 1 day after tamoxifen treatment. Again, an increase in the frequency of tdTom+ crypts was observed (Figures 1P–1R). Following dextran sodium sulfate (DSS)-induced colitis, multicrypt tdTom+ patches (MCPs) were detected at the margins of regions of damage (Figures 1S–1U;Figures S1T and S1U). Together, these results suggest that Atoh1+ cells directly contribute to regeneration following damage.

**Creating a Pro-secretory Phosphomutant ATOH1**

Previous studies have indicated that multisite phosphorylation of bHLH proteins restrains cell cycle exit and limits differentiation, whereas, conversely, un(der)phosphorylation promotes these processes in the developing nervous system and pancreas (Ali et al., 2011, 2014; Azzarelli et al., 2017). However, a role for multisite phospho-regulation of bHLH proteins in adult homeostasis or tissue repair has not been reported. Hence, we hypothesized a potential role for ATOH1 phosphorylation in controlling the transition between stem and progenitor compartments both in homeostasis and under conditions of heightened proliferation following tissue damage. Cyclin-dependent kinases phosphorylate on serine-proline (SP) or threonine-proline (TP) residues. ATOH1 has 9 S/T-P sites available for phosphorylation (Figures 2A–2C). ATOH1 can be phosphorylated on many sites; we observed at least 5 distinct phospho-forms of ATOH1 after phosphorylation by different Cyclin and Cdk combinations (Figure 2B). We expressed forms of ATOH1 where S/T-P sites were mutated.
to alanine-proline (AP) in colorectal cancer cells to determine the effect of ATOH1 phosphorylation on cell proliferation and on expression of markers of differentiation. The phosphorylation of two SP sites has previously been shown to destabilize the ATOH1 protein in the context of neuronal precursors (Forget et al., 2014). Although mutation of these two phospho-sites had a modest effect on ATOH1 activity, mutation of all 9 S/T-P sites was more effective at promoting enhanced cell cycle exit (Figures 2D and 2E). Additionally, the expression of secretory genes (Figure 2F) was enhanced after mutation of all 9 potential phosphorylation sites compared with both wild-type ATOH1, 2S-A, ATOH1, and 7S/T-A ATOH1. These observations are consistent with multisite phospho-regulation of ATOH1 playing a significant role in controlling the balance between proliferation and differentiation, as described for other bHLH family members (Ali et al., 2011, 2014; Azzarelli et al., 2017).

**9S/T-A Phosphomutant ATOH1 Promotes Secretory Maturation In Vivo**

To investigate how preventing phosphorylation of ATOH1 affects progenitor-mediated self-renewal in homeostasis and repair, we substituted 9S-T-A ATOH1 for the wild-type form in its endogenous locus, generating a knockin mouse identical to Atoh1(WT)CreERT2, but with the hyperactive phosphomutant Atoh1(9S/T-A)CreERT2 allele (Figure S2A). Homozygous Atoh1(9S/T-A)CreERT2 and control Atoh1(WT)CreERT2 mice were generated. Phenotype analysis identified no gross differences between the two lines. Mice developed normally, and the overall morphological appearance of the epithelium remained unchanged. More detailed analysis found no difference in the number or distribution of the different secretory lineages or in the frequency of apopotic cells (Figures S2B–S2F).

To investigate whether the 9S/T-A mutations affect secretory maturation after lineage specification, transcriptional profiling of secretory cells in the two lines was performed. First the expression profile of Atoh1+ cells from Atoh1(WT)CreERT2 mice was determined by comparing tdTom+ (secretory) and tdTom- (absorptive) cells to define the baseline pro-secretory signature for both the colon and SI (Table S1). Next, the transcription profiles of tdTom+ cells from wild-type and mutant mice (Table S2) were compared against this Atoh1+ baseline and a published secretory signature (Lo et al., 2016). These gene set enrichment analyses (GSEAs) demonstrated a major pro-secretory shift in Atoh1(9S/T-A)CreERT2 mice in both tissues and a strongly reduced secretory signature induced by prevention of multisite phosphorylation of Atoh1+ cells to this population, mathematical modeling was used to infer the proportion of the clonogenic fraction that is initially marked in Atoh1(WT)CreERT2 and Atoh1(9S/T-A)CreERT2 mice. In both the SI and colon, the contribution of Atoh1+ progenitors to the stem cell pool is reduced in 9S/T-A animals (Figures 3N–3Q). Between 1% and 2% of SI crypts in Atoh1(WT)CreERT2 mice contain a single clonogenic stem cell derived from an Atoh1+ progenitor, and this is reduced 5-fold in Atoh1(9S/T-A)CreERT2 mice (Figures 3N and 3Q). In the colon, values are higher, with the observed 4% wholly populated crypts (WPCs) and 5% partly populated crypts (PPCs) identified in Atoh1(WT)CreERT2 mice 30 days post-induction requiring that initially 44% of crypts (1 in 15 active stem cells) contained an Atoh1+-derived stem cell. This is reduced to 11% in 9S/T-A mutant mice (Figures 3P and 3Q). Notably, these rates reflect the contribution of a single cohort of transient progenitors arising from the stem cell pool that are produced over 1 or 2 days.

**Epithelial Proliferation and Clonogenicity Are Inhibited in Atoh1(9S/T-A) Mice**

We next investigated whether the enhanced pro-secretory signature induced by prevention of multisite phosphorylation of ATOH1 is accompanied by changes in proliferation. Comparing proliferation between the two lines demonstrated a slight overall decrease in the total proliferative index of the crypts of Atoh1(9S/T-A)CreERT2 mice in both the SI and colon, but that did not reach significance in the latter. More detailed spatial analysis within the crypt epithelium demonstrated that this effect was largely accounted for by a decrease in the proportion of cells in S phase in the epithelium of 9S/T-A mutants in cell positions above the very base of the crypt and a reduction in the frequency of proliferative goblet cells (Figures 3F–3H; Figure S3A). This supports the interpretation that the phosphorylation of ATOH1 in cells immediately arising from the stem cell population limits Atoh1+-dependent cell cycle exit to allow maintenance of proliferation in progenitors. Reciprocally, preventing this phosphorylation limits the ability to return to a proliferative stem and progenitor compartment. We next tested this hypothesis using a lineage tracing approach.

Lineage tracing and fluorescence-activated cell sorting (FACS) analysis established that the acute pattern of reporter expression and absolute number of tdTom+ cells were the same in Atoh1(9S/T-A)CreERT2 and controls (Figures S3B–S3I). However, lineage tracing at 30 days identified fewer epithelial crypts in both the SI and colon than in Atoh1(WT)CreERT2 mice (Figures 3I–3M). The 9S/T-A Atoh1+ cells were also impaired in their ability to form tdTom+ clones after radiation (Figure S3J). Together, the observations demonstrate that preventing phosphorylation of ATOH1 impairs the return of Atoh1+ cells to the stem cell compartment and confirm a role for ATOH1 phosphorylation in maintenance of progenitor plasticity.

Previously, we and others have described that only a subset of competing stem cells drive increases in clone sizes that lead to surviving clones populating entire crypts (Kozar et al., 2013; Ritsma et al., 2014). To determine the net contribution of Atoh1+ cells to this population, mathematical modeling was used to infer the proportion of the clonogenic fraction that is initially marked in Atoh1(WT)CreERT2 and Atoh1(9S/T-A)CreERT2 mice. In both the SI and colon, the contribution of Atoh1+ progenitors to the stem cell pool is reduced in 9S/T-A animals (Figures 3N–3Q). Between 1% and 2% of SI crypts in Atoh1(WT)CreERT2 mice contain a single clonogenic stem cell derived from an Atoh1+ progenitor, and this is reduced 5-fold in Atoh1(9S/T-A)CreERT2 mice (Figures 3N and 3Q). In the colon, values are higher, with the observed 4% wholly populated crypts (WPCs) and 5% partly populated crypts (PPCs) identified in Atoh1(WT)CreERT2 mice 30 days post-induction requiring that initially 44% of crypts (1 in 15 active stem cells) contained an Atoh1+-derived stem cell. This is reduced to 11% in 9S/T-A mutant mice (Figures 3P and 3Q). Notably, these rates reflect the contribution of a single cohort of transient progenitors arising from the stem cell pool that are produced over 1 or 2 days.

**Compromised Epithelial Regeneration in Atoh1(9S/T-A) Mice**

Although phosphorylation of ATOH1 clearly regulates reversion of secretory progenitors to the stem cell compartment, the absence of any other apparent phenotype in 9S/T-A Atoh1 mice suggests a limited requirement for such plasticity in homeostasis. We next investigated the role of ATOH1 phosphorylation in mounting a robust regenerative response following tissue damage. In the DSS-induced chemical colitis model,
9S/T-A mutant mice showed a greater sensitivity after treatment, with increased weight loss and slowed recovery (Figures 4A, S4A, and S4B). Analysis of this phenotype at the start of the regenerative phase (9 days after the start of DSS treatment) showed areas of ulceration that were more extensive in mice carrying the 9S/T-A mutant (Figures 4B and 4C). At both 5 and 9 days, the proportion of secretory cells was identical for the two lines, and cell death was restricted to a few cells on the luminal surface, suggesting that the greater sensitivity does not arise from enhanced damage or deletion of secretory cells in 9S/T-A mutant mice (Figures 4D–4F). However, lineage tracing 30 days following DSS treatment identified a reduced number and size of tdTom+ regenerative patches in 9S/T-A colons compared with the wild-type (WT) (Figures 4G, 4H, S4C, and S4D). Together, these results demonstrate that mice lacking the ability to phosho-regulate ATOH1 have compromised regenerative capacity following damage and that the contribution of Atoh1+ progenitors is required for robust tissue repair.

DISCUSSION

It is now accepted that cells with the capacity for self-renewal arise from a larger population whose members all have the same self-renewal potential subject to occupying available niches (Farin et al., 2016; Ritsma et al., 2014). Here we show that Atoh1+ cells make a more substantial contribution to stem cell maintenance from cells committing to secretory differentiation than has been recognized so far (van Es et al., 2012;
Self-renewal is therefore not solely a feature driven from a fixed pool of stem cells but, rather, involves dynamic interchange between progenitors and stem cells in the steady state. Transcription factors of the bHLH family have been extensively studied as master regulators of cell fate commitment and differentiation in a wide variety of tissues, including the nervous system and intestine (Ali et al., 2011, 2014; Yang et al., 2001). However, in recent years, additional roles for these proteins are emerging in direct coordination of cell cycle and differentiation events, particularly during embryonic development (Castro and Guillemot, 2011). Intestinal homeostasis in many ways represents an ongoing development-like hierarchical process where crypts are maintained by stem cells feeding a proliferating progenitor compartment that gives rise to a variety of mature cell types. What is now also emerging is a picture of significant plasticity where cells expressing Atoh1, previously thought to represent a population that has undergone secretory commitment, can nevertheless revert to “stemness” and repopulate the entire crypt with surprisingly high frequency. The mechanisms controlling this plasticity have been unclear. Here we determine that the balance between stem and progenitor fate behavior in the intestine can be controlled by Atoh1 multisite phosphorylation under normal homeostatic conditions.

Control of proliferation and differentiation by modulation of bHLH protein phosphorylation is emerging as an important mechanism in development of the nervous system and the pancreas (Cleaver, 2017; Guillemot and Hassan, 2017). We now demonstrate that multisite phosphorylation is also required to restrain irreversible commitment of secretory precursors in the adult homeostatic gut and so to maintain their ability to repopulate the stem cell compartment. Consistent with this, a phosphomutant form of ATOH1 enhances the expression of gene sets associated with a more mature secretory phenotype in colorectal carcinoma cells. Interestingly, in the homeostatic gut, despite Atoh1+ cells normally supplying up to 1 in 15 cells in the stem cell compartment, the phosphomutant Atoh1-expressing intestine is essentially phenotypically normal, indicating that plasticity from the secretory to the stem compartment is not essential in normal homeostasis. However, intestinal regeneration after damage is substantially compromised by an inability to phosphorylate ATOH1.
Taken together, our results indicate that multisite phosphorylation of ATOH1 is used to dynamically regulate the return of secretory precursors to the stem cell compartment, which facilitates the capacity of the epithelium as a whole to respond rapidly to changes in the cellular environment. Damaging the intestine using irradiation or DSS (van Es et al., 2012; Ishibashi et al., 2018) leads to acute cell damage and death, followed by proliferative regeneration that produces new cells for tissue repair. Activation of cyclin-dependent kinases (CDKs) and mitogen-activated protein kinases (MAPKs) in rapidly proliferating cells undergoing regeneration would result in enhanced phosphorylation of ATOH1, restraining further progression down the secretory lineage and supporting re-entry of Atoh1-expressing cells into a stem-like state. The post-translational regulation of ATOH1 by proline-directed kinases to modulate the balance between proliferation and differentiation in response to changing tissue demands in the adult intestinal epithelium echoes the regulation and effect of other bHLH proteins as development progresses (Hardwick et al., 2015).

The secretory fate choice mediated by ATOH1, a master regulator, is not irreversible differentiation; rather, it is entry into a plastic state through which progression is regulated by post-translational modifications. Functionally, the implications are likely to be that post-translational modifications facilitate rapid cellular responses by allowing reversal of commitment or varying its extent or rate. Progenitor plasticity is not merely an incidental acquired behavior following damage but plays an integral part in tissue restoration and requires post-translational regulation of ATOH1.

STAR METHODS

Detailed methods are provided in the online version of this paper and include the following:

- **KEY RESOURCES TABLE**
- **CONTACT FOR REAGENT AND RESOURCE SHARING**
- **EXPERIMENTAL MODEL AND SUBJECT DETAILS**
  - Mice
  - Cell Lines
- **METHOD DETAILS**
  - Cloning of mouse knock-in constructs
  - ES cell targeting
  - Mouse genotyping
  - Creation of doxycycline inducible DLD-1 cells
  - Treatment of animals
  - Crypt fractionation and single cell preparation
  - Flow cytometry
  - Whole-mount preparation
  - Whole-mount scanning and quantification
  - Antibody staining
  - Quantification of crypts in whole-mounts
  - Immunostainings
  - Single molecule FISH
  - Analysis of gut sections
  - Colon ulceration scoring
  - RNA isolation
  - Gene expression analysis
  - RNA sequencing
  - Western blotting
  - In vitro kinase assay
  - Cell proliferation and cell cycle analysis
  - RNA sequencing analysis
  - Secretory signature gene list
  - Gene Set Enrichment Analysis (GSEA)
  - QUANTIFICATION AND STATISTICAL ANALYSIS
    - Computational analysis
    - Model fitting
    - Statistical analysis
  - DATA AND SOFTWARE AVAILABILITY

SUPPLEMENTAL INFORMATION

Supplemental Information includes four figures and three tables and can be found with this article online at https://doi.org/10.1016/j.stem.2018.07.002.

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AUTHOR CONTRIBUTIONS

Conceptualization, D.J.W., A.P., and R.K.; Methodology, G.T., D.J.W., and A.P.; Formal Analysis, G.T., E.M., and C.S.R.C.; Investigation, G.T., S.K., S.B.-M., A.H., and R.A.; Writing – Original Draft, D.J.W., A.P., and G.T.; Writing – Review & Editing, G.T., D.J.W., and A.P.; Visualization, G.T.; Supervision, D.J.W., A.P., and S.I.; Project Administration, D.J.W. and A.P.; Funding Acquisition, D.J.W. and A.P.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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# STAR METHODS

## KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Antibodies**      |        |            |
| Mouse Monoclonal anti-Atoh1 | Developmental Studies Hybridoma Bank | Cat# Math1 (Atoh1): RRID:AB_10805299 |
| Rabbit Polyclonal anti-β-actin | Abcam | Cat# ab8227: RRID:AB_2305186 |
| IRDye 800CW Goat anti-Mouse IgG (H + L) | LI-COR Biosciences | Cat# P/N 925-32210: RRID:AB_2687825 |
| IRDye 680LT Goat anti-Rabbit IgG (H + L) | LI-COR Biosciences | Cat# P/N 925-68021: RRID:AB_2713919 |
| Rat Anti-Mouse CD326 (Ep-CAM) Monoclonal Antibody, Alexa Fluor 488 Conjugated, Clone G8.8 | BioLegend | Cat# 118210: RRID:AB_1134099 |
| Sheep Polyclonal BrdU Antibody | Abcam | Cat# ab1893: RRID:AB_302659 |
| Rabbit Polyclonal Anti Human Lysozyme | Dako | Cat# A0099: RRID:AB_2341230 |
| Biotin-SP-AffiniPure Donkey Anti-Sheep IgG (H+L) | Jackson ImmunoResearch Labs | Cat# 713-066-147: RRID:AB_2340717 |
| Biotin-SP-AffiniPure Donkey Anti-Rabbit IgG (H+L) | Jackson ImmunoResearch Labs | Cat# 711-065-152: RRID:AB_2340593 |
| Rabbit Anti-Chromogranin A Polyclonal Antibody | Abcam | Cat# ab15160: RRID:AB_301704 |
| Rabbit Polyclonal Anti-Synaptophysin Antibody | Millipore | Cat# AB9272: RRID:AB_570874 |
| Rabbit Polyclonal anti-DCAMKL1 Antibody | Abcam | Cat# ab31704: RRID:AB_873537 |
| Rabbit Anti-Human Lysozyme Polyclonal Antibody, FITC Conjugated | Dako | Cat# F037021: RRID:AB_578661 |
| Rabbit Anti-Mucin 2 Polyclonal Antibody | Santa Cruz Biotechnology | Cat# sc-15334: RRID:AB_2146667 |
| Donkey anti-Rabbit IgG (H+L) Secondary Antibody, Alexa Fluor 488 | Thermo Fisher Scientific | Cat# A-21206: RRID:AB_2534792 |

| **Bacterial and Virus Strains** |        |            |
|---------------------------------|--------|------------|
| RP24-77K22 Bacterial Artificial Chromosome | BACPAC Resources Center | N/A |

| **Chemicals, Peptides, and Recombinant Proteins** |        |            |
|---------------------------------------------------|--------|------------|
| Lambda Protein Phosphatase (Lambda PP) | New England Biolabs | Cat# P0753S |
| Doxycycline Hydrochloride, Ready Made Solution | Sigma-Aldrich | Cat# D3072 |
| Tet Approved FBS | Clontech Laboratories | Cat# 631101 |
| Tamoxifen | Sigma-Aldrich | Cat# T5648 |
| Dextran Sulfate Sodium Salt | MP Biomedicals | Cat# 02160110 |

| **Critical Commercial Assays** |        |            |
|--------------------------------|--------|------------|
| In-Fusion HD Cloning Kit | Clontech Laboratories | Cat# 639648 |
| TruSeq Stranded mRNA Library Prep Kit | Illumina | Cat# 20020595 |

| **Deposited Data** |        |            |
|--------------------|--------|------------|
| RNA sequencing data | This paper | GEO: GSE115416 |
| Mendeley Data | This paper | https://doi.org/10.17632/vgydv5b949.1 |

| **Experimental Models: Cell Lines** |        |            |
|------------------------------------|--------|------------|
| DLD-1 Flip-In T-Rex cell line | Laboratory of Stephen Taylor | N/A |

| **Experimental Models: Organisms/Strains** |        |            |
|-------------------------------------------|--------|------------|
| Atoh1^{WT}/CreER{T2} | This paper | N/A |
| Atoh1^{9S/T-A}/CreER{T2} | This paper | N/A |

(Continued on next page)
CONTACT FOR REAGENT AND RESOURCE SHARING

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Douglas J. Winton (doug.winton@cruk.cam.ac.uk).

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Mice

Mice used in this study were 8-16 weeks old males and females of C57BL/6 background. The mice were housed under controlled conditions (temperature (21 ± 2 °C), humidity (55 ± 10%), 12 h light/dark cycle) in a specific-pathogen-free (SPF) facility (tested according to the recommendations for health monitoring by the Federation of European Laboratory Animal Science Associations). The animals had unrestricted access to food and water, were not involved in any previous procedures and were test naive. All experiments were carried out on homozygous Atoh1(WT)CreERT2 and Atoh1(9S/T-A)CreERT2 lines. For lineage tracing experiments, the mice were heterozygous for the reporter gene (Rosa26tdTom/+). All animal experiments were carried out in accord with the guidelines of the UK Home Office, under the authority of a Home Office project license approved by the Animal Welfare and Ethical Review Body at the CRUK Cambridge Institute, University of Cambridge.

Cell Lines

DLD-1 (human colon adenocarcinoma, male) cells, modified with the Flp-In T-Rex system (Thermo Fisher), were used in the study. The cell line authentication was carried out using Single Tandem Repeat (STR) genotyping. Tests were performed routinely to confirm mycoplasma-negative status of the cells. The cells were maintained in Dulbecco’s Modified Eagle Medium (DMEM) with L-glutamine. Medium was supplemented with 10% Tet System-approved fetal bovine serum (FBS, Clontech). The cells were cultured under standard conditions (5% CO₂, 37 °C).
METHOD DETAILS

Cloning of mouse knock-in constructs
For generation of mouse knock-ins Atoh1 locus and homology arms were amplified from a bacterial artificial chromosome (BAC) RP24-77K22 (BACPAC Resources Centre). The targeting construct was assembled by a combination of seamless cloning (In-Fusion, Clontech) and restriction digest and ligation. For this a loxP site was introduced into 5′UTR of Atoh1 via PCR amplification. A neomycin cassette was inserted such that the 3′UTR was not disrupted. The CreERT2-hCD2-3′UTR was generated via gene synthesis service (Integrated DNA Technologies). The Atoh1 sequence (Atoh1WT or Atoh1RM57-A9) was merged with this construct, and then ligated with Atoh1 vector containing the homology arms. The targeting vector sequence was verified by Sanger sequencing and linearized by Swal enzyme before transfecting into ES cells. The final inserted sequence is available on request.

ES cell targeting
Electroporation of the targeting construct into mouse ES cells was conducted by the CRUK CI Transgenic Core. ES cells were positively selected with G418. Correct integration of the construct was verified by long range PCR (SequlalPrep, Thermo Fisher) according to the manufacturer’s instructions. Left integration arm was detected using a forward primer 5′-GGA CAG GCG GGA ACC ACA GA-3′ and a reverse primer 5′-TTG TCA ACA CGA GCT GGT CGA A-3′. Right integration arm was amplified using the following set of primers: forward 5′- CAA CAC AAC CCT GAC CTG TG-3′, and reverse 5′- CCC TAA CCA GTG TGC CCT TA-3′. Copy number of the clones was determined by qPCR of the neomycin selection cassette via a commercial genotyping service provider (Transnetyx). Single copy ES cell clones were taken forward for blastocyst injection, and chimeric mice were generated. Following successful germline transmission, the mice heterozygous for the targeting construct were crossed onto PGK-Cre line (Lallemand et al., 1998) in order to remove both the neo selection cassette and the endogenous Atoh1 locus at the same time. A constitutively active Atoh1-P2A-CreERT2 allele was generated in this process.

Mouse genotyping
Genotyping was carried out by Transnetyx. Manual genotyping by PCR was used to distinguish between homozygous and heterozygous Atoh1 animals. The following primers were used: forward 5′-TTT GTT GTT GTT CGG GG-3′; reverse 5′-TCT TTT ACC TCA GCC CAC TCT T-3′.

Creation of doxycycline inducible DLD-1 cells
To generate an inducible stable cell line, a DLD-1 Flp-In T-Rex cell line containing a single Frt site was obtained (a generous gift from Prof Stephen Taylor, University of Manchester). Atoh1 construct in a pcDNA 5/FRT/TO vector (Thermo Fisher) was co-transfected to generate an inducible stable cell line, a DLD-1 Flp-In T-Rex cell line containing a single Frt site was obtained (a generous gift from Prof Stephen Taylor, University of Manchester). Atoh1 construct in a pcDNA 5/FRT/TO vector (Thermo Fisher) was co-transfected to generate an inducible stable cell line, a DLD-1 Flp-In T-Rex cell line containing a single Frt site was obtained (a generous gift from Prof Stephen Taylor, University of Manchester). Atoh1 construct in a pcDNA 5/FRT/TO vector (Thermo Fisher) was co-transfected

Treatment of animals
Induction of CreERT2 in animals was carried out using the free base tamoxifen (Sigma) dissolved in ethanol/oil (1:9). The animals received 3 mg tamoxifen via an intra-peritoneal injection in all experiments. To define Atoh1 secretory signature, the mice were injected with 1 mg tamoxifen per day on 3 consecutive days for maximal labeling of all secretory lineages.

SI injury was induced by exposing animals to whole-body irradiation (6 Gy). To induce colon-specific injury, mice were given 1.5% DSS (MP Biomedicals) in drinking water for 5 days. DSS was replaced every two days during the treatment. To induce lineage tracing and ablate Lgr5+ cells in Lgr5DTR mice, the animals first received 3 mg tamoxifen i.p., followed by an i.p. injection of DT in saline (50 µg/kg) 6 h later.

Crypt fractionation and single cell preparation
Sl (proximal 15 cm) and colon were dissected, flushed with PBS, everted and fed onto a glass rod spiral. They were incubated at 37°C in Hank’s Balanced Salt Solution (HBSS) without Ca2+ and Mg2+, containing 10 µM EDTA and 10 mM NaOH. Crypt release was facilitated using a vibrating stirrer (Chemap). Samples were incubated for 1 h and pulsed every 10 min. Fractions were collected after each pulse, and fresh solution added. Crypt-enriched fractions were pooled and washed in cold 2% FBS/PBS. Fraction 1 (villus-enriched)
was discarded. Pooled fractions were resuspended in 0.05% trypsin and incubated for 7 min at 37°C, shaking every 1 min. Single cells were then filtered through a 70 μm mesh, and washed twice in 2% FBS/PBS.

Flow cytometry
Single cell suspension obtained by trypsin treatment was washed and incubated with an anti-mouse CD326 (EpCAM) AlexaFluor 647 antibody (1:2,000, clone G8.8, Biolegend). DAPI (10 μg/mL) was added to distinguish between live and dead cells. Flow sorting was carried out on a BD FACS Aria SORP (BD Biosciences), using appropriate single-stained and unstained controls.

Whole-mount preparation
Tissue was cut open, pinned out luminal side up, and fixed for 3 h at room temperature in ice-cold 4% PFA in PBS (pH 7.4). Whole-mounts were washed with PBS, and incubated with demucifying solution (3 mg/mL dithiothreitol (DTT), 20% ethanol, 10% glycerol, 0.6% NaCl, 10 mM Tris, pH 8.2) for 20 min, and mucus removed by washing with PBS.

Whole-mount scanning and quantification
The tdTom fluorescence in colon whole-mounts was detected using Amersham Typhoon 5 laser scanner (GE Healthcare) at a 10 μm resolution. The tdTom+ foci were scored manually in Fiji. Mid and distal colon were scored only as the shape of the proximal colon prevented confident assessment of tdTom+ patches.

Antibody staining
For staining whole-mount sections of 2 cm in length were excised, washed in 0.1% PBS-T for 2 days, and blocked in 10% donkey serum in PBS overnight at 4°C, protected from light. Samples were then incubated with an anti-mouse CD326 (EpCAM) AlexaFluor 647 antibody (1:100, clone G8.8, Biolegend) in 10% donkey serum in PBS for 3 days. Finally, the tissue was washed with PBS-T for 1 day.

Quantification of crypts in whole-mounts
Imaging was done on a TCS SP5 confocal microscope (Leica). Images were analyzed using Fiji. For SI, a minimum of 2,500 crypts per animal was scored. For colon, at least 900 crypts per mouse were scored. For the low-power analysis of clonal events, tdTom+ clones were scored across the entire length of the SI whole-mounts using a stereomicroscope (Nikon).

Immunostainings
For immunohistochemistry SI and colon were opened and fixed for 24 h in 10% neutral buffered formaldehyde in PBS. The tissue was paraffin embedded and sectioned by the CRUK CI Histopathology core. Haematoxylin and eosin staining was performed using an automated ST5020 Multistainer (Leica Biosystems). Alcian Blue and Periodic Acid/Schiff staining was carried out by the CI Histopathology Core. Briefly, slides were incubated in Alcian Blue for 10 min, and washed in water. They were then incubated in 0.5% periodic acid for 5 min, and washed 3 times. Slides were incubated in Schiff’s reagent for 15 min, washed 3 times, and counterstained with Mayer’s Haematoxylin.

BrdU and lysozyme immunohistochemistry was carried out using a Bond Max autostainer (Leica), with a proteinase K antigen retrieval. Slides were blocked with 3% hydrogen peroxide, followed by incubation in Avidin/Biotin Blocking Kit (Vector Laboratories). BrDU was detected using a sheep anti-BrdU antibody (1:500, Abcam ab1893). Rabbit anti-lysozyme antibody (1:500, Dako A0099) was used for lysozyme staining. Secondary antibodies in the two cases were biotinylated donkey anti-sheep (1:250, Jackson ImmunoResearch 713-066-147) and biotinylated donkey anti-rabbit (1:250, Jackson ImmunoResearch 711-065-152), respectively. Slides were incubated with Streptavidin coupled with horseradish peroxidase (HRP), and color developed using diaminobenzidine (DAB) and DAB Enhancer (Leica).

Synaptophysin and Chromogranin A detection was carried out by manual IHC. Antigen retrieval was performed with 10 mM citrate buffer (pH 6.0) in a pressurised heating chamber. Tissue sections were incubated with rabbit anti-Chromogranin A antibody (1:500, Abcam ab15160), rabbit anti-Synaptophysin antibody (1:300, Millipore AB9272), overnight at 4°C. Slides were incubated with biotinylated donkey anti-rabbit secondary antibody (1:500, Jackson ImmunoResearch 711-065-152), Streptavidin-HRP conjugate (Vector Laboratories) was added onto the slides and incubated for 30 min. DAB Chromogen substrate (Dako) was added for dye development. Counterstaining and dehydration was performed on the ST5020 Multistainer (Leica) followed by coverslipping.

For immunofluorescence tissue was excised and fixed for 48 h in 4% PFA in PBS at 4°C, after which it was transferred to 20% sucrose solution. After cryosectioning antigen retrieval where needed was accomplished by incubating the slides in 1% SDS for 5 min. Blocking was performed with 5% donkey serum. Following a wash primary antibodies were added and incubated overnight at 4°C. The following primary antibodies were used: rabbit FITC-anti-Lyz (1:400, Dako, F037201), rabbit anti-Muc2 (1:50, Santa Cruz, sc-15334), rabbit anti-ChgA (1:100, Abcam, ab15160), and rabbit anti-Dclk1 antibody (1:1000, Abcam, ab31704). Secondary detection was with AlexaFluor 488 donkey anti-rabbit secondary antibody (1:500, Thermo Fisher, A-21206). Alkaline phosphatase activity was detected using Blue AP kit (Vector Laboratories). Sections were covered with Prolong Gold with DAPI (Life Technologies). Fluorescent imaging was carried out on a TCS SP5 confocal microscope (Leica).
**Single molecule FISH**

Harvested SI and colon tissues were flushed with cold 4% formaldehyde (FA) in PBS and incubated first in 4% FA/PBS for 3 hours, then in 30% sucrose in 4% FA/PBS overnight at 4°C with constant agitation. Fixed tissues were embedded in OCT. Quantification of co-expression was achieved by smFISH. Probe library design, hybridization procedures, and imaging settings were carried out according to published methods (Itzkovitz et al., 2011; Lyubimova et al., 2013). A Nikon-Ti-E inverted fluorescence microscope equipped with a Photometrics Pixis 1024 CCD camera was used to image a 10 μm cryo-section. A stack of 30 frames with 0.3 μm intervals was acquired to allow 3D cell imaging. FITC-conjugated antibody for E-cadherin was added to the hybridization mix and used to visualize cell borders. Detection of cells that were positive for Lgr5 transcripts, Atoh1 transcripts or both was performed manually with Fiji.

**Analysis of gut sections**

Stained longitudinal sections of the SI and colon were visualized and positive cells scored manually. BrdU+ and negative nuclei were scored in complete half-crypt sections. Lysozyme+ cells were counted per whole crypt section. Alcian Blue and PAS+ cells were counted in complete half-villus sections, between the crypt neck and the tip of the villus. Cells in which the stain was clearly associated with a corresponding nucleus were marked as positive. Chromogranin A+ and synaptophysin+ cells were scored per complete half-crypt-villus section. Positive and negative crypts were scored, and results expressed as a frequency of positive cells.

**Colon ulceration scoring**

H&E-stained sections of colons were scanned on Aperio slide scanner (Leica Biosystems), and analyzed using eSlide Manager (Leica Biosystems). Ulceration was defined as a region of a complete loss of crypt architecture and high cellularity.

**RNA isolation**

For gene expression analysis by qPCR, cells were lysed and RNA isolated using RNeasy Mini Plus kit (QIAGEN). For sequencing, total RNA was isolated from flow-sorted cells using RNeasy Micro Plus kit (QIAGEN).

**Gene expression analysis**

RNA was converted into cDNA (iScript cDNA synthesis kit, BioRad), and gene expression was analyzed using TaqMan gene expression probes (Thermo Fisher). The following probes were used: Atoh1 (Mm00476035_s1), Muc2 (Hs00894053_g1), Tff3 (Hs00902278_m1), Spdef (Hs01026050_m1), Dll4 (Hs00184092_m1), Rassf4 (Hs00604698_m1), Gapdh (Hs02758991_g1). All TaqMan assays are listed in Table S3.

**RNA sequencing**

Samples for RNA sequencing were collected 24 h post-tamoxifen induction (3 mg i.p. injection). The tissue was fractionated as described above and cells prepared for flow cytometry. The cells were stained and sorted in the same way as for other experiments, as noted above. EpCAM+ttdTomlive cells were collected directly into the lysis buffer and RNA was extracted immediately following the sort (RNeasy Micro Plus Kit, QIAGEN). RNA quality was assessed on a 2100 Bioanalyser instrument (Agilent), according to the manufacturer’s instructions. The libraries were prepared using TruSeq Stranded mRNA Library Prep Kit (Illumina) and sequenced as 50 bp single-end reads on the Illumina HiSeq 4000 system.

**Western blotting**

Protein extracts for SDS-PAGE were prepared by lysing the cells with RIPA buffer containing protease and phosphatase inhibitor cocktail (Thermo Fisher). Mouse anti-ATOH1 antibody (1:100, Developmental Studies Hybridoma Bank) and a rabbit anti-β-A antibody (1:5,000, ab8227, Abcam) were used. Fluorescent secondary antibodies were used (Li-Cor, goat anti-mouse 800LT (1:5,000), goat anti-rabbit 680LT (1:20,000)). For some experiments, protein extracts were incubated with λ phosphatase (New England Biolabs) prior to western blotting, according to the manufacturer’s instructions.

**In vitro kinase assay**

The assay was performed as previously described (Azzarelli et al., 2017), with minor modifications. HA-tagged WT and mutant ATOH1 were in vitro translated (TNT® Quick Coupled Transcription/Translation Systems, Promega) in the presence of LiCl (800 mM) to reduce potential phosphorylation in reticulocyte lysate. Samples were incubated with human recombinant CDK/Cyclins (0.25 μM final concentration) in the presence of 10 μM ATP for 1 h at 30°C. Proteins were separated on Phos-tag gels (Alpha Laboratories, 7.5% acrylamide, 50 μM phos-tag PAGE, Wako) and immunoblotted with rat anti-HA-Peroxidase (1:5000, Roche).

**Cell proliferation and cell cycle analysis**

Cell proliferation was assessed by an automated live-cell imaging system (IncuCyte ZOOM, Essen Bioscience). For cell cycle analysis, the cells were trypsinised, washed, fixed with ethanol, and stained with propidium iodide prior to flow cytometry.
RNA sequencing analysis
The reads were aligned to the mouse reference genome [GRCm38] using TopHat2 aligner (Kim et al., 2013). Differentially expressed gene lists were generated using DESeq2 package from Bioconductor (Love et al., 2014).

Secretory signature gene list
The list of differentially expressed genes (p < 0.01) was generated by comparing the transcripts from tdTom+ and tdTom− cells of Atoh1(WT)CreERT2 Rosa26tdTom/+ mice following tamoxifen. Upregulated genes in tdTom+ cells were selected to define a secretory signature in the small intestine and colon (Table S1). The top 500 upregulated, differentially expressed genes were used to perform the Gene Set Enrichment Analysis (GSEA).

Gene Set Enrichment Analysis (GSEA)
This analysis was performed using the GSEA software from the Broad Institute (http://software.broadinstitute.org/gsea/index.jsp) (Subramanian et al., 2005). The list comprised all differentially expressed and non-differentially expressed genes from the 9S/T-A v WT comparison in SI and colon, respectively. This gene list was probed with the previously generated secretory signatures (top 500 upregulated genes), and the published Atoh1+ gene signatures for ileum, colon (Lo et al., 2016), and intestinal stem cells (Muñoz et al., 2012).

QUANTIFICATION AND STATISTICAL ANALYSIS
Computational analysis
The process by which crypt stem cells replace each occurs in a random though predictable manner. This behavior can be modeled via a stochastic birth-death process (Lopez-Garcia et al., 2010; Snippert et al., 2010). The model was derived to model experiments where a single stem cell is labeled in a handful of crypts. As the number of initially labeled crypts was not of interest and to bypass any variability coming from the initial induction, the different time points comparable, the equations were rescaled to account for only the surviving clones. Here we know the parameters of the stem cell dynamics (Kozar et al., 2013; Vermeulen et al., 2013), and would like to know the starting number of labeled stem cells per crypt and the number of labeled crypts.

For this analysis we use the equations described previously (Lopez-Garcia et al., 2010; Snippert et al., 2010), reproduced below. The probability of a crypt having clone of size n (for 0 < n < N) at time t is:

\[ p_n(t) = \frac{2}{N} \sum_{m=1}^{N-1} (-1)^{m+n+1} \cos\left(\frac{\pi m}{2N}\right) \left(1 - e^{-\lambda t} \sin^2\left(\frac{\pi m}{2N}\right) t\right) \] (Equation 1)

Here n is the number of labeled stem cells, N is the total number of stem cells, \( \lambda \) is the rate of stem cell replacement. And for the probability of all stem cells labeled we have:

\[ p_N(t) = \frac{2}{N} \sum_{m=1}^{N-1} \left(1 - e^{-\lambda t} \sin^2\left(\frac{\pi m}{2N}\right) t\right) \] (Equation 2)

These equations assume the initial conditions of one labeled stem cell at t = 0. The starting labeled stem cells were chosen randomly at the beginning of each simulation.

The values we observe for the clonal frequencies are substantially lower than what the model would predict, suggesting that not all crypts have labeled stem cells. In order to find out the fraction of labeled crypts \( \nu \) we use a mixture model:

\[ Q_n(t) = (1 - \nu) \delta_0 + \nu P_n(t) \] (Equation 3)

Where \( Q_n(t) \) is the probability that a randomly selected crypt has a clone of size n labeled stem cells at time t. We use the values of N, \( \lambda \) and \( \nu \) from Kozar et al. (2013) and Vermeulen et al. (2013) and estimate \( \nu \).

Model fitting
For every mouse, at day 30 we count the number of clones (\( k_i \)) and the number of crypts (\( C_i )\). We use a hierarchical model to capture the mouse to mouse variability. The statistical model is a follows

\[ k_i \sim\text{Binomial}(C_i, R_i(30 - \tau)) \] (Equation 4)

\[ R_i(30 - \tau) \sim\text{Student}_t(\eta, Q(30 - \tau), \sigma) \] (Equation 5)

Here \( R_i \) is truncated to [0, 1]. For the SI no distinction is made in clone size, so Q is the sum of all \( Q_n \) and for the colon we use only the full clones for fitting \( Q = Q_N \).

The priors on the population parameters are:

\[ \eta \sim\text{Gamma}(2.0.1) \] (Equation 6)
\[ \sigma \sim \text{Gamma}(0.01, 0.01) \]  
(Equation 7)

The prior on the mixing coefficient is

\[ \eta \sim \text{Beta}(1/2, 1/2) \]  
(Equation 8)

The posterior was derived via MCMC using Rstan (Carpenter et al., 2017). For the proximal and distal SI we used \( \tau = 5 \) as the clones were measured in ribbons coming out of the crypt, which take a few days to emerge from the crypt base. Whereas for the colon we used \( \tau = 1 \). The parameters used were \( \tau = 0.1, N = 5 \) for proximal SI, \( \tau = 0.2, N = 6 \) for distal and \( \tau = 0.3, N = 7 \) for colon.

**Statistical analysis**

Statistical tests were not used to predetermine sample size. Randomization was not performed to allocate samples/animals to experimental groups. Blinding was performed for quantifications in Figures 3F and 3G, as well as Figures S2B–S2F. Data analysis was performed using GraphPad Prism software or R package.

**DATA AND SOFTWARE AVAILABILITY**

The accession number for the RNA sequencing data reported in this paper is GEO: GSE115416. Mendeley Dataset of original data can be accessed at https://doi.org/10.17632/vgvdv5b949.1.