



Natural Selection, Crypt Fitness, and Pol III Dependency in the Intestine

The Harvard community has made this
article openly available. [Please share](#) how
this access benefits you. Your story matters

Citation	Jadhav, Unmesh, and Ramesh A. Shivdasani. 2016. "Natural Selection, Crypt Fitness, and Pol III Dependency in the Intestine." Cellular and Molecular Gastroenterology and Hepatology 2 (6): 714-715. doi:10.1016/j.jcmgh.2016.09.006. http://dx.doi.org/10.1016/j.jcmgh.2016.09.006 .
Published Version	doi:10.1016/j.jcmgh.2016.09.006
Citable link	http://nrs.harvard.edu/urn-3:HUL.InstRepos:31731662
Terms of Use	This article was downloaded from Harvard University's DASH repository, and is made available under the terms and conditions applicable to Other Posted Material, as set forth at http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#LAA

EDITORIAL

Natural Selection, Crypt Fitness, and Pol III Dependency in the Intestine



The intestinal epithelium is a rapidly self-renewing tissue. The enormous replicative burden is borne by stem cells, which in mice divide approximately once a day, and by the transit-amplifying compartment, where cells replicate 2–3 times more frequently. Because the continual production of many new cells creates substantial demands, the intestine of conditional mutant mice is an ideal organ to examine tissue-specific functions of pervasive housekeeping factors.

Beyond the sheer pace of self-renewal, the intestine offers other advantages. Arguably no other site shows the relationship of mother and daughter cells—to each other and to the surrounding niche—as clearly. In addition, lineage tracing through cell-restricted expression of Cre recombinase allows monitoring of cells' immediate and distant progeny, an approach that has contributed much to the current understanding of stem and progenitor cells. Finally, the intestinal epithelium is exceptionally adaptive. Intact crypts compensate rapidly for defective ones, first by increasing the rate of cell proliferation and then by undergoing fission to produce new crypts; this fission resembles the process of intestinal growth in fetuses and children. The gut mucosa is thus a Darwinian terrain: as a population, crypts continually express a fitness that is necessary to maintain vital barrier and absorptive functions. In this issue of *Cellular and Molecular Gastroenterology and Hepatology*, Kieckhafer et al¹ exploit tissue-specific gene disruption and the experimental assets of the mouse intestine to identify the cells most vulnerable to the absence of a “housekeeping” factor.

The DNA-dependent enzyme RNA polymerase III (Pol III) synthesizes noncoding transfer RNAs (tRNAs) and other transcripts associated with RNA splicing and protein synthesis. Its highest activity occurs during cell division and organogenesis, and the enzyme is essential for organismal viability. Pol III subunit B (Polr3B), the second largest protein in the complex, is important for enzyme structure and function, and human *POLR3B* mutations cause neurologic defects and leukodystrophy rooted in hypomyelination.² In mice, loss of the Pol III repressor MAF1 increases precursor tRNAs in many tissues and the resulting increase in energy expenditure may underlie resistance to obesity.³ A hypomorphic mutation in the zebrafish *Polr3b* gene, *slim jim*, provided early clues that optimal Pol III activity is necessary for replication of intestinal progenitor cells,⁴ and Kieckhafer et al¹ reported the consequences of a similar tissue-restricted mutation in the mouse gut epithelium.

tRNA levels were reduced, as expected, and *Polr3b* mutant mice grew slower than their litter mates, with a minority surviving beyond the first week of life. These

findings reflect both a profound deficit in cell proliferation and increased death of mutant crypt cells. The associated morphologic defects were absent in embryos or neonates, even though the *Villin-Cre* transgene the investigators used to disrupt *Polr3b* becomes active well before the onset of embryonic villus morphogenesis.⁵ Thus, the critical need for Pol III activity in the intestine seems to be surmounted during development and becomes overt only after birth, during a period when brisk cell replication is necessary to support organ growth as well as new nutritional and absorptive demands.

Kieckhafer et al¹ also observed many crypts that contained an abundance of proliferating cells, showed ongoing Wnt pathway activity, lacked dying cells, and produced normal villi. Cells in these intact crypts had escaped Cre-mediated recombination of one or both *Polr3b* gene copies, and over time they occupied larger fractions of the mucosa. This process of natural selection recapitulates many other instances of loss of genes linked to vital cellular processes, such as mitosis and modulation of chromatin.⁶ The presence and expansion of these adaptive crypts enables intestinal function in surviving animals.

The findings reported by Kieckhafer et al¹ thus highlight the acute need for protein synthesis to sustain perinatal intestinal growth and function, as well as the tissue's remarkable versatility in the face of crypt attrition. The strain of *Polr3b* mutant mice reported in this study will help determine how other tissues respond, at various developmental stages and under stressful conditions, to reduced levels of Pol III-dependent tRNAs and other transcripts. With regard to the intestine, 2 questions come to mind. The demand for protein synthesis and cell replication seems, on the surface, at least as high in the fetal as in the neonatal gut. One now wonders if this is indeed true, and, if so, how the fetal intestine withstands the paucity of vital RNAs. More generally, it is important to understand how intact crypts sense and respond to defects in their neighbors, because that understanding holds one key to improved treatments for inflammatory and other bowel disorders.

UNMESH JADHAV, PhD

RAMESH A. SHIVDASANI, MD, PhD

Department of Medical Oncology and Center for Functional Cancer Epigenetics

Dana-Farber Cancer Institute

Department of Medicine

Brigham and Women's Hospital

Harvard Medical School

Boston, Massachusetts

References

1. Kieckhafer JE, Lukovac S, Ye DZ, et al. The RNA Polymerase III Subunit Polr3b is required for the maintenance of small intestinal crypts in mice. *Cell Mol Gastroenterol Hepatol* 2016;2:783–795.
2. Bernard G, Vanderver A. Pol III-related leukodystrophies. In: Pagon RA, Adam MP, Ardinger HH, et al., eds. *GeneReviews*. Seattle, WA: University of Washington, 1993.
3. Bonhoure N, Byrnes A, Moir RD, et al. Loss of the RNA polymerase III repressor MAF1 confers obesity resistance. *Genes Dev* 2015;29:934–947.
4. Yee NS, Gong W, Huang Y, et al. Mutation of RNA Pol III subunit *rpc2/polr3b* leads to deficiency of subunit *Rpc11* and disrupts zebrafish digestive development. *PLoS Biol* 2007;5:e312.
5. Madison BB, Dunbar L, Qiao XT, et al. Cis elements of the villin gene control expression in restricted domains of the vertical (crypt) and horizontal (duodenum, cecum) axes of the intestine. *J Biol Chem* 2002;277:33275–33283.
6. Jadhav U, Nalapareddy K, Saxena M, et al. Acquired tissue-specific promoter bivalency is a basis for PRC2 necessity in adult cells. *Cell* 2016;165:1389–1400.

Correspondence

Address correspondence to: Ramesh A. Shivdasani, MD, PhD, Dana-Farber Cancer Institute, Department of Medical Oncology, 450 Brookline Avenue, Boston, Massachusetts 02215. e-mail: ramesh_shivdasani@dfci.harvard.edu.

Conflicts of interest

The authors disclose no conflicts.

Funding

Supported by F32DK103453 (U.J.) and R01DK081113 (R.A.S.).

Most current article

© 2016 The Authors. Published by Elsevier Inc. on behalf of the AGA Institute. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

2352-345X

<http://dx.doi.org/10.1016/j.jcmgh.2016.09.006>