
Review

How to Heal the Gut's Brain: Regeneration of the Enteric Nervous System

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Abstract: The neural-crest derived enteric nervous system (ENS) is the intrinsic nervous system of the gastrointestinal (GI) tract and controls all gut functions, including motility. Lack of ENS neurons causes various ENS disorders such as Hirschsprung Disease. One treatment option for ENS disorders includes the activation of resident stem cells to regenerate ENS neurons. Regeneration in the ENS has mainly been studied in mammalian species using surgical or chemically-induced injury methods. These mammalian studies showed a variety of regenerative responses with generally limited regeneration of ENS neurons, but (partial) regrowth and functional recovery of nerve fibers. Several aspects might contribute to the variety in regenerative responses, including observation time after injury, species, and gut region targeted. Zebrafish have recently emerged as a promising model system to study ENS regeneration as larvae possess the ability to generate new neurons after ablation. As the next steps in ENS regeneration research, we need a detailed understanding of how regeneration is regulated on a cellular and molecular level both in animal models with high and low regenerative capacity. Understanding the regulatory programs necessary for robust ENS regeneration will pave the way for using neural regeneration as a therapeutic approach to treating ENS disorders.

Keywords: enteric progenitor cell; zebrafish; inflammation; Hirschsprung Disease; neural crest cell; ENS neuropathies

1. Introduction

The enteric nervous system (ENS) is derived from the neural crest cell lineage and provides the intrinsic innervation of the gastrointestinal (GI) tract forming a complex network of different types of neurons and glial cells [1-3]. It is the largest division of the peripheral nervous system – the ENS contains approximately the same number of neurons as the adult spinal cord [4-6]. The mammalian ENS consists of two enteric plexi, the myenteric plexus, and the submucosal plexus. ENS neurons are found in ganglia, which are connected by a network of nerve fibers [5]. As the intrinsic nervous system of the gut, the ENS regulates many essential intestinal functions such as GI motility, absorption of nutrients, secretion, fluid exchange, regulation of blood flow, epithelial barrier function, immune modulation, and microbiota colonization and composition [5, 7-9]. The ENS regulates intestinal functions along the entire length of the gut - in mammals, the GI tract can be subdivided into upper and lower parts. The upper GI tract contains the stomach and small intestine (duodenum, jejunum, and ileum), the lower GI tract the cecum, colon, and rectum (**Figure 1**).

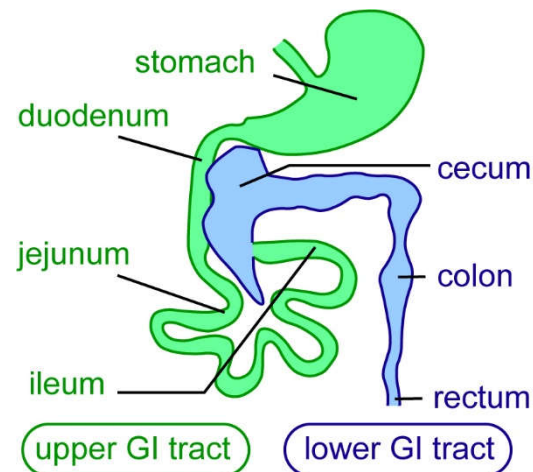


Figure 1. Subdivisions of the mammalian gastrointestinal tract. In mammals, the gastrointestinal (GI) tract, here exemplified in mice, is divided into an upper (green) and a lower GI tract (blue). The upper GI tract consists of duodenum, jejunum, and ileum. The lower GI tract consists of the cecum, colon, and rectum. A. modified from [10].

Each of these gut regions is exposed to daily abrasive and potentially harmful and toxic compounds which in turn subject the ENS to many stressors and mechanical forces. As the ENS directly overlays the intestinal smooth muscle layer it changes its shape during the gastrointestinal contractions and relaxations that accompany intestinal motility [11]. In aging animals, ENS neuron numbers are considerably reduced [11-13]. In addition, acute and chronic gut disorders can also impact ENS cells. Deficits in ENS neuron abundance and composition cause severe GI dysfunction that occur in congenital ENS disorders such as Hirschsprung disease, inflammatory gut diseases like inflammatory bowel syndrome, and neurodegenerative diseases, such as Parkinson's Disease [14-19]. ENS defects in these various disorders range from a complete deficit of neurons in a gut subdivision to a lack of specific neuronal subtypes. For example, in Hirschsprung Disease, ENS neurons are lacking in the distal gut resulting in difficulties in passing stool. The length of the aganglionic part varies from distal colon to the entire colon and stretches of the small intestine [15].

1.1. Therapeutic approaches to treat ENS disorders

At present, ENS disorders are treated symptomatically or require surgical removal of the area with ENS neuron deficits [15, 18, 20, 21]. Because of the prevalence of ENS disorders and their strong impact on the patient's quality of life, there has been an increasing interest in finding therapeutic approaches for restoring lost ENS neurons or glial cells.

There are two main avenues for treating such ENS disorders: ENS stem-cell-based treatment or stimulation of resident stem cells to regenerate missing ENS cells [20]. Stem-cell-based therapeutic approaches aim to transplant ENS stem cells into a patient's gut where they then can differentiate into ENS neurons and/or ENS glia cells [20, 22]. In mammals including humans, enteric neuronal stem cell (ENSC) populations are present into adulthood [22, 23]. When isolated and induced in culture, ENSCs can be amplified and differentiated into many enteric neuronal and glial subtypes [22, 24]. Additionally, recent work has generated different types of ENS neurons and glia cells from induced pluripotent stem cells (iPSCs) [25, 26]. Transplantation of ENSCs in mouse models of congenital ENS disorders shows their ability to migrate and colonize mammalian guts [22, 24, 27]. Yet, so far, such stem cell-

based therapeutic approaches have not been translated into the clinic and practical limitations remain [27]. As this review focuses on ENS regeneration, an in-depth discussion of stem cell-based therapies is out of its scope. Additional information on ENSCs, their origins, *in vitro/in vivo* experiments, and their therapeutic potential can be found in several excellent recent reviews [15, 24, 27-29].

The other avenue for treating insufficient or injured ENS neurons in patients is to stimulate resident stem cells to regenerate the missing/damaged enteric neurons. For this approach to be successful, a detailed understanding of the extent, potential, and cellular and molecular mechanisms underlying ENS regeneration is necessary. In this review, we will discuss the ability of the ENS to regenerate and why ENS regeneration might be limited in mammals. We will also discuss the high regenerative capacity of the ENS in the zebrafish model system and close with an outlook on the open questions and future directions for ENS regeneration research.

1.2. What constitutes nervous system regeneration?

Nervous system regeneration is generally defined as either the repair or the new generation of neurons damaged by injury or disease. This can occur at two different levels: First, the regrowth of just neuronal axons when the neuronal cell body is not damaged. Second, “large scale” regeneration, where new neurons are generated and have to connect to the existing neural circuitry or build a new circuit that then gets wired into the larger neural circuit [30]. Complete regeneration is viewed as the full restoration of lost neurons or neuronal function, whereas partial regeneration includes the generation of new neurons or nerve fibers without the complete restoration of lost neurons or neuronal function.

1.3. The mammalian ENS has limited regenerative ability

Starting in the 1950s, regeneration studies have been performed in the ENS in a variety of mammalian research organisms, including adult guinea pigs, rats, mice, and dogs. Very few studies have so far been performed in non-mammalian species. In mammals, neurons or nerve fibers are generated in some but not all experimental settings after injury. Generally, neuron numbers are not restored to control levels and nerve fiber regeneration is often not complete. Additionally, within the injured area, significant structural changes remain. Several aspects might contribute to this variety in the regenerative responses: type of injury, observation time after injury, animal model, and which gut region has been injured.

In the next section, we will discuss the main injury models used to study ENS regeneration and how the variability in experimental parameters might impact the regenerative response. The injury models that we discuss here are surgical or mechanical methods of injury, chemical-induced injury, and infection with pathogens.

1.4. ENS regeneration after surgical/mechanical injury in mammals

Transection and reanastomosis and (partial) stenosis are the main types of surgical injury that have been used to study ENS regeneration in various mammalian animal models (**Figure 2**).

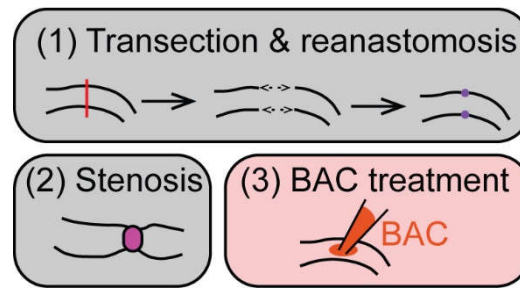


Figure 2. Common ENS injury approaches in mammals. Common models using surgical (grey boxes) or chemically-induced (red) injury methods to study ENS regeneration in mammals: (1) transection & reanastomosis: the targeted part of the gut is transected (red line) with subsequent end-to-end anastomosis (purple). (2) For stenosis, a ring (magenta) is placed around the gut, which causes partial obstruction. (3) Benzalkonium chloride (BAC) treatment comprises the treatment of a small gut segment with BAC, which leads to a loss of ENS neurons in the treated area.

Transection and reanastomosis in the small intestine in dogs, guinea pigs, or rats resulted in nerve fiber regeneration across the injury site with accompanying functional recovery. In dogs, transection and reanastomosis led to an initial loss of migratory motor complex (MMC) propagations after surgery [31, 32]. MMCs are cyclic sweeping gut movements in the small intestine during fasting [33]. After 40-60 days post-surgery, MMCs started to be coupled again between the two segments indicating functional recovery of gut movements across the injury site. By approximately 100 days post-surgery, MMCs were fully recovered suggesting regeneration of nerve fibers across the injury site [31, 32].

Analysis of structural changes after surgical injury in the ENS in rats or guinea pigs showed that neurons degenerated in the injury site between 1-2 months after transection and reanastomosis in the small intestine [34-36]. At 6 weeks post-surgery, there were significantly fewer neurons up to about 5mm from the surgical site. Farther away, no difference in neuron numbers was seen. Even at 12 months, the surgical site did not contain any new neurons [34, 35]. The area close to the surgical site showed an increase in large extra-ganglionic neurons forming clusters, but these additional neurons did not restore neuron numbers to control levels. A little farther away, in the area that had not seen a reduction in neuron numbers, large extra-ganglionic neurons emerged at 6 months and continued to increase in numbers until reaching a plateau at 1-year post-surgery [34]. These extra-ganglionic neurons were found in higher numbers than controls, forming clusters and connecting to surrounding enteric ganglia [34]. Interestingly, in mice, extra-ganglionic neurons that move into enteric ganglia after 24 weeks were observed in the small intestine and colon after experimental treatment with a 5-HT₄ agonist to stimulate neurogenesis [37]. This indicates a potential connection between the process of adult neurogenesis and regeneration.

After 2-4 weeks, regenerating nerve fibers appeared in the lesion site both from the oral and aboral sides. These fibers extended further after 8 weeks, suggesting that nerve fiber regeneration across the lesion site occurs also in guinea pigs and rats [34]. Yet, structural changes in neuronal fiber patterns were visible even after 1 year post-surgery indicating that nerve fibers were not fully restored across the lesion site [34]. Together, these studies showed that transection and reanastomosis in the small intestine led to distinct structural changes in neurons and nerve fibers depending on the distance from the surgical site and time after surgery, but never fully restore structural patterns as in the uninjured gut.

Analysis of structural changes after transection and reanastomosis in the lower GI tract in guinea pigs had contrasting results [38-40]. Surgery in the colon showed substantial, and long-lasting disruption of neuron numbers and neuronal pathways at the lesion site [38]. Neurons degenerated at the injury site and failed to regenerate after 8-24 weeks. Nerve fiber regeneration occurred, but in this model, regrowth of nerve fibers occurred preferentially in an oral to anal direction, no regrowth of fibers anal to oral was observed between 10-24 weeks after surgery [38]. In contrast, transection and reanastomosis in the guinea pig rectum led to nerve fiber regrowth across the lesion site accompanied by recovery of rectal contractions between 2-8 weeks [39, 40]. In addition, new neurons were present in the injured site, but no ganglia were formed [39, 40]. The difference in neuron and nerve fiber regeneration between the different gut regions suggests that each gut subdivision may have different abilities to restore neuronal function after surgical injury.

Injury models using (partial) stenosis focused on the small intestine [41-45]. Between 1-2 weeks after stenosis in rats, there was an increase in the numbers and cell volume of neurons per ganglia accompanied by thymidine incorporation or expression of the proliferation marker PCNA in neurons upstream of the point of stenosis. This suggests that neurons either activated the cell cycle or were undergoing unscheduled DNA synthesis due to DNA repair [45]. It was not tested if the increase in neurons resulted in a full restoration of lost neurons. After 4 weeks, there was no evidence of proliferating cells, but the nuclear volume of neurons was still increased [41-43]. In a guinea pig model of stenosis, there was no evidence of an increase in neuron numbers after 3-5 weeks [44], but the earlier time points where cell cycle activation in neurons was observed in the rat models were not analyzed in the guinea pig. Thus, parallel conclusions are unable to be made.

In summary, surgical injuries result in partial nerve fiber regeneration with functional recovery, some generation of neurons, but no full recovery of neuron numbers after surgery. There is some evidence that gut regions and species differ in their regenerative ability and response, but more extensive research needs to be done in comparing the regenerative ability of different species and gut regions. Notably, regenerative processes take a long time, which is relevant for the experimental setup to study ENS regeneration.

1.5. ENS regeneration after chemically-induced injury in mammals

A chemically-induced injury method commonly used to study ENS regeneration is the treatment with benzalkonium chloride (BAC), which leads to a loss of ENS neurons in the treated area (**Figure 2**, [46, 47]). In rat or mouse animal models, treatment with BAC of the small intestine or colon resulted in neuron degeneration and denervation within 2-5 days after treatment [48-50]. In the small intestine of rats, there was a significant increase in new neurons at the lesion site at 30-60 days, which suggests a partial regenerative response with ~25-30% of control neuron numbers present 60 days post-injury [50]. In the treated mouse colon, in contrast, no consistent generation of new neurons was observed. Here, nerve fibers grew into the denervated area between 7-14 days post-treatment. Nerve fiber density increased until 60 days post-treatment to 35-60% of the control density. Occasionally new neurons were present along the nerve fiber bundles, but neuron numbers did not increase significantly [49]. Genetic lineage tracing in the small intestine in mice also found that neuronal projections grew into the injured area along with newly generated enteric glia cells, but at 3 months post-ablation, new neurons were not present within the injured area (though they were found

adjacent to the area of injury) [48]. In the rat small intestine, nerve fiber regrowth took even longer – at 15 days, no neuron fibers were present. By 45 days, some neuronal fibers had regrown but fibers showed marked structural differences to the control ENS [51].

In conclusion, injury of the ENS using chemical treatment results in denervation of the injured area. However, depending on the animal model and gut region the timing of regrowth of nerve fibers and the regenerative ability differs. As with the surgical injury method, regeneration in the colon seems to be impaired compared to the small intestine, for which some, limited regeneration of nerve fibers and ENS neurons was observed.

1.6. What aspects might impact the regenerative process in mammals?

The regeneration studies in mammals have shown that even though there are instances of nerve fiber regeneration and partial regeneration of neurons, complete regeneration with restoration of neuron numbers to control levels does not take place. In this section, we will discuss three aspects that may impact the regenerative ability in the mammalian ENS:

1.6.1. Time required for full regeneration

Regeneration events generally took a long time and positive signs of regeneration were often not observed until months after injury. The reporting period of many studies was significantly shorter and may not cover a long enough time to determine if neuron regeneration might still happen. Thus, results determining no or limited regeneration capacity may be limited by the timeframe of observation.

1.6.2. Impact of inflammation

In the mammalian central nervous system (CNS), neuroinflammation is both detrimental as well as positive for a regenerative response. In zebrafish – a research organism with a remarkable ability to regenerate nervous system injury – inflammatory responses support brain, retina, and spinal cord regeneration [52-54]. The gut contains a large number of immune cells and intestinal inflammation results in structural changes in the ENS including hyperplasia or loss of neurons [55, 56]. Inflammatory responses have been evoked by different types of treatments, for example with trinitrobenzene sulfonic acid (TSA), dextran sulfate sodium (DSS) [57, 58], or by infection with the nematode *Nippostrongylus brasiliensis* [59]. Treatment with TSA results in an increase in proliferating cells at 3 and 7d post-treatment, but the study did not investigate if this supported the generation of new neurons later [58]. DSS treatment in the colon led to a significant increase in the number of ENS neurons 2 days after treatment. In this treatment setting, a subset of enteric glia cells was suggested to give rise to ENS neurons [57, 60]. Infection with *Nippostrongylus brasiliensis* resulted in a strong inflammatory response and led to a significant degeneration of nerve fiber densities until 21 days post-infection (dpi) [59]. Reinnervation peaked at 18 dpi and experimental guts showed higher fiber density than in control animals [59]. Together, new neurons can be generated in an inflammatory setting. This suggests that inflammation might have positive effects on neural regeneration in the ENS. Even though inflammatory responses have been documented after ENS injury, many experimental settings/studies did not report if there was inflammation or not. Also, it has not directly been tested if an inflammatory response impacts the regenerative processes as it does in the CNS. Thus, further study of the role of inflammation for ENS regeneration is necessary specifically in the context of different types of injury.

1.6.3. Ability of mammalian adult neurogenesis in the ENS

In the CNS, the ability to regenerate neurons in the adult has been connected to the extent of adult neurogenesis, that is to say – high levels of adult neurogenesis correspond to the ability to regenerate the nervous system after injury. Low to no adult neurogenesis is generally correlated with low or no ability to regenerate neurons in the adult [53, 61, 62]. In the ENS, the extent of adult neurogenesis in mammals beyond postnatal stages has been debated in the field. The predominant view on neurogenesis in the mammalian ENS has been that the majority of enteric neurons are generated during embryogenesis and early postnatal stages with no continuous generation of neurons throughout adult life [43]. Only in specific experimental conditions, for example, induction of inflammation with DSS treatments or the treatment with 5-HT₄ agonists, has the generation of new neurons been observed in the adult [37, 57]. Treatment with 5-HT₄ agonists translates to improved neural regeneration, as nerve fiber regeneration and reflex recovery were accelerated in resection and anastomosis in the rectum of guinea pigs. Also, the generation of new neurons after DSS treatment was abolished by treatment with a 5-HT₄ antagonist [39, 40, 57]. This suggests that signals that support adult neurogenesis, also promote neural regeneration in the ENS. Recently, the view of little to no adult neurogenesis in the ENS has been challenged by the study of Kulkarni et al. (2016) which observed high levels of adult neurogenesis in the gut with a steady turnover of ENS neurons in homeostatic conditions [63]. Despite the lack of clarity regarding the extent of adult neurogenesis in the mammalian ENS, ENSCs can be isolated from the gut into adulthood and differentiate into ENS neurons *in vitro* [22, 24]. This shows that stem cells, which can generate new neurons are still present in the adult mammalian ENS, but work is still needed to determine how these cells can be re-activated.

1.7. The zebrafish ENS regenerates neurons after focal ablation

Recent studies have found that zebrafish generate new neurons after focal ablation of a small fraction of ENS neurons (**Figure 3A**, [64, 65]). This is consistent with the continuous generation of new ENS neurons into adulthood in zebrafish [66]. After laser-ablation of a small number of neurons in the distal zebrafish larval gut, new neurons are generated at the site of ablation, and neuron numbers increase 5 and 10 days post-ablation (dpa) at a significantly higher rate than in the control (**Figure 3B**). It remains to be determined if neuron numbers completely catch up to control levels and if these neurons restore gut function. The study by Ohno et al. (2021) also shed light on potential cellular mechanisms that drive neuronal regeneration in the zebrafish ENS (**Figure 3B**). Nerve fibers first enter the ablated area at 1 dpa forming a bridge across the ablated area. At 1 dpa, *sox10* positive enteric progenitor cells (EPCs) seem to enter the area of ablation via these nerve fiber bridges (**Figure 3B**). These EPCs proliferate indicating that progenitor cell proliferation drives ENS regeneration in zebrafish [64]. These data suggest that zebrafish can generate new neurons after injury, but the occurrence of regeneration in adult zebrafish remains to be tested. Thus, zebrafish are a promising new research model to study the cellular and molecular mechanisms underlying ENS regeneration.

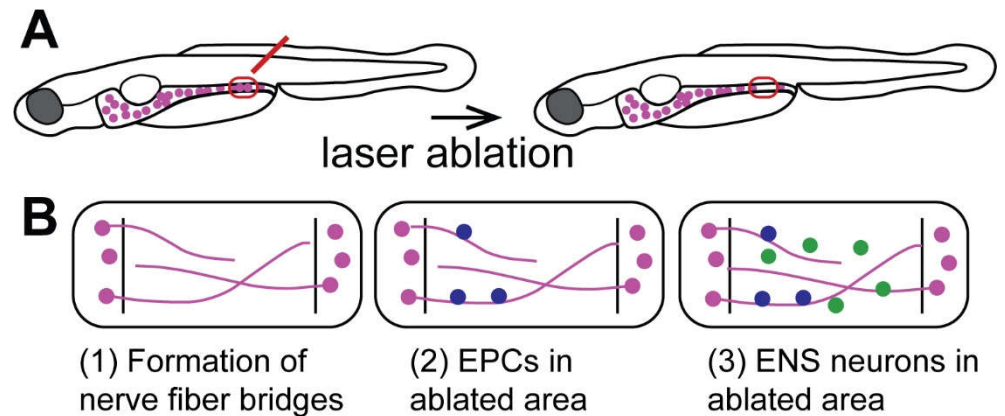


Figure 3. Regenerative processes in the zebrafish ENS after cell ablation. (A) Experimental setup of focal laser ablation (red) of a small set of ENS neurons in the zebrafish larvae. (B) Steps of regenerative processes in the zebrafish ENS in the ablated area marked by lines: (1) formation of nerve fiber bridge across the ablated area. (2) *sox10* positive enteric progenitor cells (EPCs, blue) appear in the ablated area. (3) new neurons (green) are generated in the ablated area. A-B. modified from [64].

2. Outlook

Previous studies have shown that ENS regeneration in mammals is limited. Zebrafish have been put forward as a new model organism to study ENS regeneration. The zebrafish model has the potential to answer the questions of how a robust regeneration response is controlled on a cellular and molecular level. It will be important to determine if a more extensive loss of ENS neurons still results in robust regeneration in zebrafish. Also, full functional recovery after neuron loss needs to be determined using approaches such as intestinal transit assay, or gut motility imaging [67, 68].

Important questions remain before regenerative approaches can be used to treat ENS disorders. The regenerative ability of mammalian models needs to be tested further with longer observation periods and a detailed analysis of functional recovery. This will also reveal important differences in the regenerative ability between ENS neuron types, gut regions, and species.

The field also needs a detailed understanding of how ENS regeneration is regulated on a cellular and molecular level both in animal models with robust vs. limited regenerative capacity. This will allow for the determination of 1) which cell types are present during ENS regeneration, 2) which molecular changes take place, and 3) which cell lineages generate new neurons. For example, ENS glia cells generate neurons during adult ENS neurogenesis [66, 69] and ENS glia cells have been suggested as a potential progenitor cell source in some injury settings [57, 60]. Also, the regulatory programs underlying ENS regeneration need to be determined. This information will provide the critical framework of cell biological processes and molecular-genetic factors essential for successful regeneration in the ENS. These are necessary steps for identifying differences in the cellular and molecular composition between model systems with high vs. low regenerative capacity in the ENS. Understanding these aspects of ENS regeneration will pave the way for using regenerative processes as a therapeutic approach to treating ENS disorders.

Author Contributions: Conceptualization, H.R., J.G.; Writing – Original Draft Preparation, H.R., J.G.; Writing – Review & Editing, H.R., J.G.; Supervision, J.G.; Funding Acquisition, J.G. All authors have read and agreed to the published version of the manuscript.

Funding: This research was funded by Funds from Michigan State University to J.G, by the American Neurogastroenterology and Motility Society Research Grant to J.G and by the NSF CAREER grant # 2143267 to J.G.

Conflicts of Interest: The authors declare no conflict of interest

References

1. Ganz, J., Gut feelings: Studying enteric nervous system development, function, and disease in the zebrafish model system. *Dev Dyn* **2018**, 247, (2), 268-278.
2. Heanue, T. A.; Shepherd, I. T.; Burns, A. J., Enteric nervous system development in avian and zebrafish models. *Dev Biol* **2016**, 417, (2), 129-38.
3. Rao, M.; Gershon, M. D., Enteric nervous system development: what could possibly go wrong? *Nat Rev Neurosci* **2018**, 19, (9), 552-565.
4. Belkind-Gerson, J.; Graeme-Cook, F.; Winter, H., Enteric nervous system disease and recovery, plasticity, and regeneration. *J Pediatr Gastroenterol Nutr* **2006**, 42, (4), 343-50.
5. Furness, J. B., *The enteric nervous system*. Blackwell Publishing: Oxford, 2006.
6. Furness, J. B.; Callaghan, B. P.; Rivera, L. R.; Cho, H. J., The enteric nervous system and gastrointestinal innervation: integrated local and central control. *Adv Exp Med Biol* **2014**, 817, 39-71.
7. Røig, A. S.; Mittge, E. K.; Ganz, J.; Troll, J. V.; Melancon, E.; Wiles, T. J.; Alligood, K.; Stephens, W. Z.; Eisen, J. S.; Guillemin, K., The enteric nervous system promotes intestinal health by constraining microbiota composition. *PLoS Biol* **2017**, 15, (2), e2000689.
8. Wiles, T. J.; Jemielita, M.; Baker, R. P.; Schlomann, B. H.; Logan, S. L.; Ganz, J.; Melancon, E.; Eisen, J. S.; Guillemin, K.; Parthasarathy, R., Host Gut Motility Promotes Competitive Exclusion within a Model Intestinal Microbiota. *PLoS Biol* **2016**, 14, (7), e1002517.
9. Neunlist, M.; Van Landeghem, L.; Mahe, M. M.; Derkinderen, P.; des Varannes, S. B.; Rolli-Derkinderen, M., The digestive neuronal-glia-epithelial unit: a new actor in gut health and disease. *Nat Rev Gastroenterol Hepatol* **2013**, 10, (2), 90-100.
10. Nishiyama, K.; Sugiyama, M.; Mukai, T., Adhesion Properties of Lactic Acid Bacteria on Intestinal Mucin. *Microorganisms* **2016**, 4, (3).
11. Gabella, G., On the plasticity of form and structure of enteric ganglia. *J Auton Nerv Syst* **1990**, 30 Suppl, S59-66.
12. Saffrey, M. J., Cellular changes in the enteric nervous system during ageing. *Dev Biol* **2013**, 382, (1), 344-55.
13. Saffrey, M. J., Aging of the mammalian gastrointestinal tract: a complex organ system. *Age (Dordr)* **2014**, 36, (3), 9603.
14. Brosens, E.; Burns, A. J.; Brooks, A. S.; Matera, I.; Borrego, S.; Ceccherini, I.; Tam, P. K.; Garcia-Barcelo, M. M.; Thapar, N.; Benninga, M. A.; Hofstra, R. M.; Alves, M. M., Genetics of enteric neuropathies. *Dev Biol* **2016**, 417, (2), 198-208.
15. Heuckeroth, R. O., Hirschsprung disease - integrating basic science and clinical medicine to improve outcomes. *Nat Rev Gastroenterol Hepatol* **2018**, 15, (3), 152-167.
16. Chalazonitis, A.; Rao, M., Enteric nervous system manifestations of neurodegenerative disease. *Brain Res* **2018**, 1693, (Pt B), 207-213.
17. Rao, M.; Gershon, M. D., The bowel and beyond: the enteric nervous system in neurological disorders. *Nat Rev Gastroenterol Hepatol* **2016**, 13, (9), 517-28.
18. De Giorgio, R.; Bianco, F.; Latorre, R.; Caio, G.; Clavenzani, P.; Bonora, E., Enteric neuropathies: Yesterday, Today and Tomorrow. *Adv Exp Med Biol* **2016**, 891, 123-33.
19. De Giorgio, R.; Guerrini, S.; Barbara, G.; Stanghellini, V.; De Ponti, F.; Corinaldesi, R.; Moses, P. L.; Sharkey, K. A.; Mawe, G. M., Inflammatory neuropathies of the enteric nervous system. *Gastroenterology* **2004**, 126, (7), 1872-83.
20. Burns, A. J.; Goldstein, A. M.; Newgreen, D. F.; Stamp, L.; Schafer, K. H.; Metzger, M.; Hotta, R.; Young, H. M.; Andrews, P. W.; Thapar, N.; Belkind-Gerson, J.; Bondurand, N.; Bornstein, J. C.; Chan, W. Y.; Cheah, K.; Gershon, M. D.; Heuckeroth, R. O.; Hofstra, R. M.; Just, L.; Kapur, R. P.; King, S. K.; McCann, C. J.; Nagy, N.; Ngan, E.; Obermayr, F.; Pachnis, V.; Pasricha, P. J.; Sham, M. H.; Tam, P.; Vanden Berghe, P., White paper on guidelines concerning enteric nervous system stem cell therapy for enteric neuropathies. *Dev Biol* **2016**, 417, (2), 229-51.
21. Goldstein, A. M.; Thapar, N.; Karunaratne, T. B.; De Giorgio, R., Clinical aspects of neurointestinal disease: Pathophysiology, diagnosis, and treatment. *Dev Biol* **2016**, 417, (2), 217-28.
22. Burns, A. J.; Thapar, N., Neural stem cell therapies for enteric nervous system disorders. *Nat Rev Gastroenterol Hepatol* **2014**, 11, (5), 317-28.
23. Bondurand, N.; Natarajan, D.; Thapar, N.; Atkins, C.; Pachnis, V., Neuron and glia generating progenitors of the mammalian enteric nervous system isolated from foetal and postnatal gut cultures. *Development* **2003**, 130, (25), 6387-400.
24. McCann, C. J.; Thapar, N., Enteric neural stem cell therapies for enteric neuropathies. *Neurogastroenterol Motil* **2018**, 30, (10), e13369.

25. Fattahi, F.; Steinbeck, J. A.; Kriks, S.; Tchieu, J.; Zimmer, B.; Kishinevsky, S.; Zeltner, N.; Mica, Y.; El-Nachef, W.; Zhao, H.; de Stanchina, E.; Gershon, M. D.; Grikscheit, T. C.; Chen, S.; Studer, L., Deriving human ENS lineages for cell therapy and drug discovery in Hirschsprung disease. *Nature* **2016**, 531, (7592), 105-9.
26. Barber, K.; Studer, L.; Fattahi, F., Derivation of enteric neuron lineages from human pluripotent stem cells. *Nat Protoc* **2019**, 14, (4), 1261-1279.
27. Alhawaj, A. F., Stem cell-based therapy for hirschsprung disease, do we have the guts to treat? *Gene Ther* **2021**.
28. Pan, W.; Goldstein, A. M.; Hotta, R., Opportunities for novel diagnostic and cell-based therapies for Hirschsprung disease. *J Pediatr Surg* **2021**.
29. McCann, C. J.; Borrelli, O.; Thapar, N., Stem cell therapy in severe pediatric motility disorders. *Curr Opin Pharmacol* **2018**, 43, 145-149.
30. Steward, M. M.; Sridhar, A.; Meyer, J. S., Neural regeneration. *Curr Top Microbiol Immunol* **2013**, 367, 163-91.
31. Matsumoto, T.; Sarna, S. K.; Condon, R. E.; Cowles, V. E.; Frantzides, C., Differential sensitivities of morphine and motilin to initiate migrating motor complex in isolated intestinal segments. Regeneration of intrinsic nerves. *Gastroenterology* **1986**, 90, (1), 61-7.
32. Sarna, S.; Condon, R. E.; Cowles, V., Enteric mechanisms of initiation of migrating myoelectric complexes in dogs. *Gastroenterology* **1983**, 84, (4), 814-22.
33. Deloof, E.; Janssen, P.; Depoortere, I.; Tack, J., The migrating motor complex: control mechanisms and its role in health and disease. *Nat Rev Gastroenterol Hepatol* **2012**, 9, (5), 271-85.
34. Tokui, K.; Sakanaka, M.; Kimura, S., Progressive reorganization of the myenteric plexus during one year following reanastomosis of the ileum of the guinea pig. *Cell Tissue Res* **1994**, 277, (2), 259-72.
35. Karaosmanoglu, T.; Muftuoglu, S.; Dagdeviren, A.; Durgun, B.; Aygun, B.; Ors, U., Morphological changes in the myenteric plexus of rat ileum after transection and end-to-end anastomosis. *J Anat* **1996**, 188 (Pt 2), 323-31.
36. Galligan, J. J.; Furness, J. B.; Costa, M., Migration of the myoelectric complex after interruption of the myenteric plexus: intestinal transection and regeneration of enteric nerves in the guinea pig. *Gastroenterology* **1989**, 97, (5), 1135-46.
37. Liu, M. T.; Kuan, Y. H.; Wang, J.; Hen, R.; Gershon, M. D., 5-HT₄ receptor-mediated neuroprotection and neurogenesis in the enteric nervous system of adult mice. *J Neurosci* **2009**, 29, (31), 9683-99.
38. Brookes, S. J.; Lam, T. C.; Lubowski, D. Z.; Costa, M.; King, D. W., Regeneration of nerve fibres across a colonic anastomosis in the guinea-pig. *J Gastroenterol Hepatol* **1996**, 11, (4), 325-34.
39. Matsuyoshi, H.; Kuniyasu, H.; Okumura, M.; Misawa, H.; Katsui, R.; Zhang, G. X.; Obata, K.; Takaki, M., A 5-HT₄-receptor activation-induced neural plasticity enhances in vivo reconstructs of enteric nerve circuit insult. *Neurogastroenterol Motil* **2010**, 22, (7), 806-13, e226.
40. Katsui, R.; Kojima, Y.; Kuniyasu, H.; Shimizu, J.; Koyama, F.; Fujii, H.; Nakajima, Y.; Takaki, M., A new possibility for repairing the anal dysfunction by promoting regeneration of the reflex pathways in the enteric nervous system. *Am J Physiol Gastrointest Liver Physiol* **2008**, 294, (4), G1084-93.
41. Benninghoff, V. o. n. A., Vermehrung und Vergrößerung von Nervenzellen bei Hypertrophie des Innervationsgebietes*. *Z. Naturforsch.* **1951**, 6b, 38-41.
42. Jew, J. Y.; Williams, T. H.; Gabella, G.; Zhang, M. Q., The intestine as a model for neuronal plasticity. *Arch Histol Cytol* **1989**, 52 Suppl, 167-80.
43. Joseph, N. M.; He, S.; Quintana, E.; Kim, Y. G.; Nunez, G.; Morrison, S. J., Enteric glia are multipotent in culture but primarily form glia in the adult rodent gut. *J Clin Invest* **2011**, 121, (9), 3398-411.
44. Gabella, G., Size of neurons and glial cells in the intramural ganglia of the hypertrophic intestine of the guinea-pig. *J Neurocytol* **1984**, 13, (1), 73-84.
45. Corvetti, G.; Fornaro, M.; Geuna, S.; Poncino, A.; Giacobini-Robecchi, M. G., Unscheduled DNA synthesis in rat adult myenteric neurons: an immunohistochemical study. *Neuroreport* **2001**, 12, (10), 2165-8.
46. Sato, A.; Yamamoto, M.; Imamura, K.; Kashiki, Y.; Kunieda, T.; Sakata, K., Pathophysiology of aganglionic colon and anorectum: an experimental study on aganglionosis produced by a new method in the rat. *J Pediatr Surg* **1978**, 13, (4), 399-435.
47. Fox, D. A.; Epstein, M. L.; Bass, P., Surfactants selectively ablate enteric neurons of the rat jejunum. *J Pharmacol Exp Ther* **1983**, 227, (2), 538-44.
48. Laranjeira, C.; Sandgren, K.; Kessaris, N.; Richardson, W.; Potocnik, A.; Vanden Berghe, P.; Pachnis, V., Glial cells in the mouse enteric nervous system can undergo neurogenesis in response to injury. *J Clin Invest* **2011**, 121, (9), 3412-24.
49. Hanani, M.; Ledder, O.; Yutkin, V.; Abu-Dalu, R.; Huang, T. Y.; Hartig, W.; Vannucchi, M. G.; Faussone-Pellegrini, M. S., Regeneration of myenteric plexus in the mouse colon after experimental denervation with benzalkonium chloride. *J Comp Neurol* **2003**, 462, (3), 315-27.
50. Ramalho, F. S.; Santos, G. C.; Ramalho, L. N.; Kajiwar, J. K.; Zucoloto, S., Myenteric neuron number after acute and chronic denervation of the proximal jejunum induced by benzalkonium chloride. *Neurosci Lett* **1993**, 163, (1), 74-6.
51. Luck, M. S.; Dahl, J. L.; Boyeson, M. G.; Bass, P., Neuroplasticity in the smooth muscle of the myenterically and extrinsically denervated rat jejunum. *Cell Tissue Res* **1993**, 271, (2), 363-74.

52. Kyritsis, N.; Kizil, C.; Brand, M., Neuroinflammation and central nervous system regeneration in vertebrates. *Trends Cell Biol* **2014**, 24, (2), 128-35.
53. Bosak, V.; Murata, K.; Bludau, O.; Brand, M., Role of the immune response in initiating central nervous system regeneration in vertebrates: learning from the fish. *Int J Dev Biol* **2018**, 62, (6-7-8), 403-417.
54. Nagashima, M.; Hitchcock, P. F., Inflammation Regulates the Multi-Step Process of Retinal Regeneration in Zebrafish. *Cells* **2021**, 10, (4).
55. Linden, D. R.; Couvrette, J. M.; Ciolino, A.; McQuoid, C.; Blaszyk, H.; Sharkey, K. A.; Mawe, G. M., Indiscriminate loss of myenteric neurones in the TNBS-inflamed guinea-pig distal colon. *Neurogastroenterol Motil* **2005**, 17, (5), 751-60.
56. Vasina, V.; Barbara, G.; Talamonti, L.; Stanghellini, V.; Corinaldesi, R.; Tonini, M.; De Ponti, F.; De Giorgio, R., Enteric neuroplasticity evoked by inflammation. *Auton Neurosci* **2006**, 126-127, 264-72.
57. Belkind-Gerson, J.; Hotta, R.; Nagy, N.; Thomas, A. R.; Graham, H.; Cheng, L.; Solorzano, J.; Nguyen, D.; Kamionek, M.; Dietrich, J.; Cherayil, B. J.; Goldstein, A. M., Colitis induces enteric neurogenesis through a 5-HT4-dependent mechanism. *Inflamm Bowel Dis* **2015**, 21, (4), 870-8.
58. Bradley, J. S., Jr.; Parr, E. J.; Sharkey, K. A., Effects of inflammation on cell proliferation in the myenteric plexus of the guinea-pig ileum. *Cell Tissue Res* **1997**, 289, (3), 455-61.
59. Stead, R. H.; Kosecka-Janiszewska, U.; Oestreicher, A. B.; Dixon, M. F.; Bienenstock, J., Remodeling of B-50 (GAP-43)- and NSE-immunoreactive mucosal nerves in the intestines of rats infected with *Nippostrongylus brasiliensis*. *J Neurosci* **1991**, 11, (12), 3809-21.
60. Belkind-Gerson, J.; Graham, H. K.; Reynolds, J.; Hotta, R.; Nagy, N.; Cheng, L.; Kamionek, M.; Shi, H. N.; Aherne, C. M.; Goldstein, A. M., Colitis promotes neuronal differentiation of Sox2+ and PLP1+ enteric cells. *Sci Rep* **2017**, 7, (1), 2525.
61. Kaslin, J.; Ganz, J.; Brand, M., Proliferation, neurogenesis and regeneration in the non-mammalian vertebrate brain. *Philos Trans R Soc Lond B Biol Sci* **2008**, 363, (1489), 101-22.
62. Grandel, H.; Brand, M., Comparative aspects of adult neural stem cell activity in vertebrates. *Dev Genes Evol* **2013**, 223, (1-2), 131-47.
63. Kulkarni, S.; Micci, M. A.; Leser, J.; Shin, C.; Tang, S. C.; Fu, Y. Y.; Liu, L.; Li, Q.; Saha, M.; Li, C.; Enikolopov, G.; Becker, L.; Rakhilin, N.; Anderson, M.; Shen, X.; Dong, X.; Butte, M. J.; Song, H.; Southard-Smith, E. M.; Kapur, R. P.; Bogunovic, M.; Pasricha, P. J., Adult enteric nervous system in health is maintained by a dynamic balance between neuronal apoptosis and neurogenesis. *Proc Natl Acad Sci U S A* **2017**, 114, (18), E3709-E3718.
64. Ohno, M.; Nikaïdo, M.; Horiuchi, N.; Kawakami, K.; Hatta, K., The enteric nervous system in zebrafish larvae can regenerate via migration into the ablated area and proliferation of neural crest-derived cells. *Development* **2021**, 148, (2).
65. El-Nachef, W. N.; Bronner, M. E., De novo enteric neurogenesis in post-embryonic zebrafish from Schwann cell precursors rather than resident cell types. *Development* **2020**, 147, (13).
66. McCallum, S.; Obata, Y.; Fourli, E.; Boeing, S.; Peddie, C. J.; Xu, Q.; Horswell, S.; Kelsh, R.; Collinson, L.; Wilkinson, D.; Pin, C.; Pachnis, V.; Heanue, T. A., Enteric glia as a source of neural progenitors in adult zebrafish. *Elife* **2020**, 9.
67. Ganz, J.; Baker, R. P.; Hamilton, M. K.; Melancon, E.; Diba, P.; Eisen, J. S.; Parthasarathy, R., Image velocimetry and spectral analysis enable quantitative characterization of larval zebrafish gut motility. *Neurogastroenterol Motil* **2018**, e13351.
68. Field, H. A.; Kelley, K. A.; Martell, L.; Goldstein, A. M.; Serluca, F. C., Analysis of gastrointestinal physiology using a novel intestinal transit assay in zebrafish. *Neurogastroenterol Motil* **2009**, 21, (3), 304-12.
69. Boesmans, W.; Lasrado, R.; Vanden Berghe, P.; Pachnis, V., Heterogeneity and phenotypic plasticity of glial cells in the mammalian enteric nervous system. *Glia* **2014**.