

## The enteric nervous system and neurogastroenterology

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**Abstract** | Neurogastroenterology is defined as neurology of the gastrointestinal tract, liver, gallbladder and pancreas and encompasses control of digestion through the enteric nervous system (ENS), the central nervous system (CNS) and integrative centers in sympathetic ganglia. This Review provides a broad overview of the field of neurogastroenterology, with a focus on the roles of the ENS in the control of the musculature of the gastrointestinal tract and transmucosal fluid movement. Digestion is controlled through the integration of multiple signals from the ENS and CNS; neural signals also pass between distinct gut regions to coordinate digestive activity. Moreover, neural and endocrine control of digestion is closely coordinated. Interestingly, the extent to which the ENS or CNS controls digestion differs considerably along the digestive tract. The importance of the ENS is emphasized by the life-threatening effects of certain ENS neuropathies, including Hirschsprung disease and Chagas disease. Other ENS disorders, such as esophageal achalasia and gastroparesis, cause varying degrees of dysfunction. The neurons in enteric reflex pathways use a wide range of chemical messengers that signal through an even wider range of receptors. These receptors provide many actual and potential targets for modifying digestive function.

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### Introduction

The gastrointestinal tract differs from all other peripheral organs in that it has an extensive intrinsic nervous system, termed the enteric nervous system (ENS) that can control functions of the intestine even when it is completely separated from the central nervous system (CNS).<sup>1</sup> The ENS, however, is not autonomous. Indeed, neuronal control of gastrointestinal function is an integrated system involving interactions between local enteric reflexes, reflexes that pass through sympathetic ganglia and reflexes that pass from the gut and back through the CNS (Figure 1). Conventional textbook descriptions of the autonomic nervous system depict efferent pathways from the CNS as two neurons in series, a preganglionic and a postganglionic neuron, and depict sensory information flowing from the periphery to the CNS through spinal and cranial primary afferent neurons. The organization of the ENS and of neuronal pathways to and from the intestine does not, however, follow these conventional concepts of the organization of the nervous system (Figure 1). For example, axons of neurons with cell bodies in the ENS (called intestinofugal neurons) project to sympathetic ganglia, the pancreas, gallbladder and trachea, and to the spinal cord and brain stem (Figure 1).<sup>2</sup>

The ENS is composed of small aggregations of nerve cells, enteric ganglia, the neural connections between these ganglia, and nerve fibers that supply effector tissues, including the muscle of the gut wall, the epithelial lining, intrinsic blood vessels and gastroenteropancreatic

endocrine cells (Figure 2). The total number of enteric neurons in humans is 400–600 million, which is greater than the total of all sympathetic and parasympathetic ganglia combined and approximately equal to the number of neurons in the spinal cord.<sup>2</sup> Figure 2 is representative of the ENS of the mammalian small intestine; the differences in structure and organization between regions of the gastrointestinal tract and species have been reviewed elsewhere.<sup>2–4</sup> Differences include the absence of a ganglionated submucosal plexus in the esophagus and stomach (organs that lack the large fluid fluxes across the mucosal epithelium that occur in the small and large intestines), the presence of a diffuse network of ganglia in the pancreas and a single layer of ganglia in the intestinal submucosa of small mammals. This Review is confined to discussion of monogastric mammals, in which most investigations have been done and which are arguably most relevant to humans.

### Importance of the ENS

The ENS has multiple roles: determining the patterns of movement of the gastrointestinal tract; controlling gastric acid secretion; regulating movement of fluid across the lining epithelium; changing local blood flow; modifying nutrient handling; and interacting with the immune and endocrine systems of the gut.<sup>2</sup> The ENS also contributes, along with glial cells, to maintaining the integrity of the epithelial barrier between the gut lumen and cells and tissues within the gut wall.<sup>5,6</sup> The importance of the ENS is highlighted by the wide range of enteric neuropathies that are caused following a failure

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### Competing interests

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in one or more of its roles. These neuropathies have been grouped as congenital or developmental neuropathies; sporadic and acquired neuropathies; neuropathies associated with other disease states (which can be secondary to the other disease); and iatrogenic or drug-induced neuropathies (Box 1).

In Hirschsprung disease, the ganglia of the ENS fail to develop in the distal bowel, but all other tissue components are intact and functional.<sup>7</sup> Under these circumstances, no propulsive activity occurs in the aganglionic bowel, and the neonate will die if this region is not removed. Degeneration of colonic enteric neurons in Chagas disease, which is precipitated by infection with the protozoan *Trypanosoma cruzi*, causes colorectal propulsion to fail and megacolon to develop in adults—similar to the problems associated with Hirschsprung disease in children.<sup>8</sup> An underlying occult tumor (that is, paraneoplastic syndrome) can also cause gut propulsion to fail, often with evidence of bowel dilatation.<sup>9,10</sup> Other enteric neuropathies that have severe effects on the motor functions of the digestive tract include esophageal achalasia, gastroparesis, hypertrophic pyloric stenosis and intestinal pseudo-obstruction.<sup>11</sup>

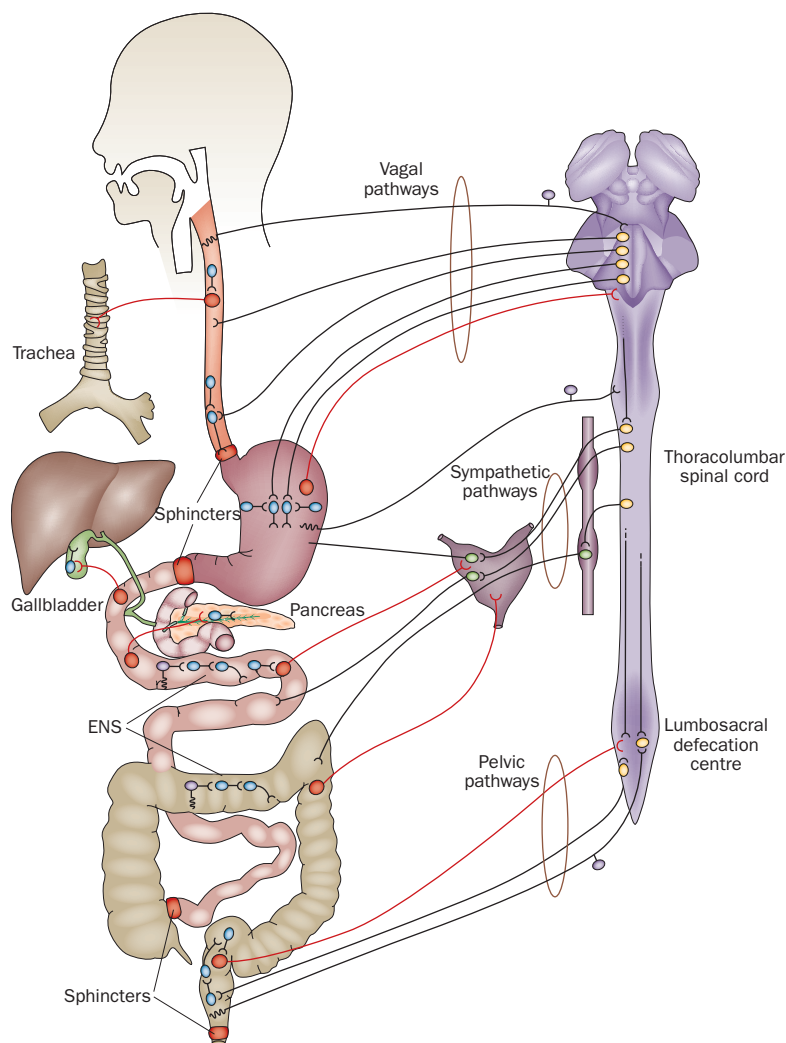
The control of fluid movement between the intestinal lumen and body fluid compartments, as discussed below, is also subject to pathological, life-threatening, influences. Fluid movement is influenced by enteric secretomotor neurons that are abnormally activated by certain infective agents or their products. These pathogens, including *Vibrio cholerae* (which secretes cholera toxin) and rotavirus (which secretes the enterotoxin NSP4), act directly on secretomotor neurons and mucosal epithelial cells,<sup>12</sup> triggering hypersecretion and subsequent diarrhea. Infectious diarrhea causes approximately 1.5 million deaths a year, primarily in underdeveloped tropical countries.<sup>13</sup>

### Control of gastrointestinal muscles

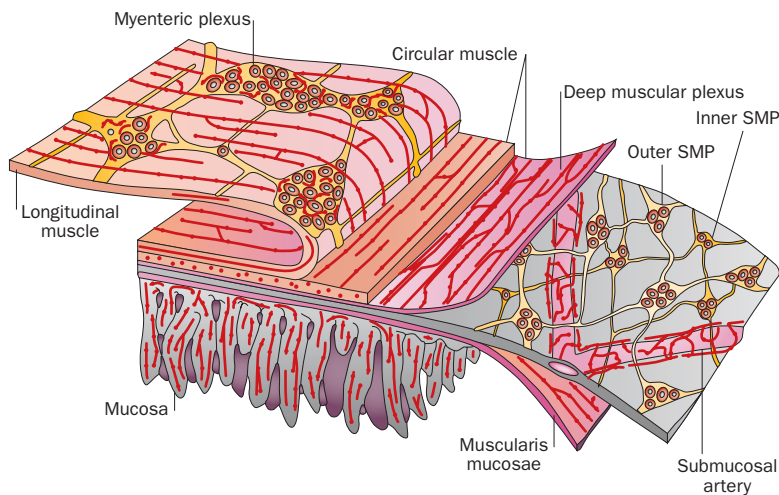
The muscle layers of the gastrointestinal tract direct propulsion, mixing of contents, reservoir capacity (notably in the stomach) and expulsion of pathogens and noxious chemicals. The degree to which the ENS is essential for coordinated muscle function, and the extent to which nerve pathways that originate outside the alimentary tract are necessary for adequate control, varies with the region of the gastrointestinal tract and also with the physiological circumstance. In broad terms, movements of the esophagus are largely determined by neural pattern generators in the CNS (that is, brain-stem circuits located in the medulla oblongata), whereas the rather extensive ENS of the esophagus has a subsidiary role. Gastric propulsive movements are primarily myogenic, but the CNS, through the brain stem and esophagogastric and gastrogastric reflexes, has a major role in monitoring the state of the stomach and, in turn, controlling its volume and strength of contractions, as well as acid secretion.<sup>14,15</sup> By contrast, the ENS dominates the control of the motility of the small and large intestines,<sup>2</sup> with the exception of defecation, for which the CNS has control through the defecation centers in the lumbosacral spinal cord.<sup>16</sup>

### Key points

- The enteric nervous system (ENS) is an extensive reflex control system for digestive function that works with the central nervous system (CNS) and neural pathways that pass through sympathetic ganglia
- The ENS of the small intestine and colon has complete reflex pathways that control patterns of contractile activity, local blood flow and transmucosal movement of fluids
- The CNS has essential roles in control of esophageal, stomach and colorectal functions
- Control of transmucosal fluid movement by the ENS and CNS is closely integrated
- The ENS interacts with both the gut endocrine and immune systems and has roles in modifying nutrient absorption and maintaining the mucosal barrier
- Enteric neuropathies in which control of muscle contractile activity or neural control of transmucosal fluid movement fail are life-threatening



**Figure 1** | The innervation of the gastrointestinal tract. The neural connections between the ENS and CNS, and neural connections between gastrointestinal organs, are quite different from those depicted in textbooks. The digestive system contains full reflex circuits of the ENS (motor neurons and interneurons in blue, sensory neurons in purple). Pathways from the gastrointestinal tract project outwards, via intestinofugal neurons (red), to the CNS (neurons in yellow), sympathetic ganglia, gallbladder and pancreas. Neurons in sympathetic prevertebral ganglia (green) receive both CNS and ENS inputs. Sensory information goes both to the ENS, via intrinsic primary afferent (sensory) neurons (purple) and to the CNS via extrinsic primary afferent neurons (also purple) that follow spinal and vagal afferent routes. Pathways from the CNS reach the ENS and gastrointestinal effector tissues through vagal, sympathetic and pelvic pathways. Abbreviations: CNS, central nervous system; ENS, enteric nervous system.



**Figure 2** | The organization of the ENS of human and medium–large mammals. The ENS has ganglionated plexuses, the myenteric plexus between the longitudinal and circular layers of the external musculature and the SMP that has outer and inner components. Nerve fiber bundles connect the ganglia and also form plexuses that innervate the longitudinal muscle, circular muscle, muscularis mucosae, intrinsic arteries and the mucosa. Innervation of gastroenteropancreatic endocrine cells and gut-associated lymphoid tissue is also present, which is not illustrated here. Abbreviations: ENS, enteric nervous system; SMP, submucosal plexus. Modified with permission from Furness, J. B. *The Enteric Nervous System* (Blackwell, Oxford, 2006).

**Box 1** | A classification of enteric neuropathies

**Congenital or developmental neuropathies**

- Hirschsprung disease (colorectal aganglionosis)
- Hypertrophic pyloric stenosis
- Multiple endocrine neoplasia 2B
- Neuronal intestinal dysplasia
- Mitochondriopathies affecting enteric neurons

**Sporadic and acquired neuropathies**

- Chagas disease
- Neurogenic forms of intestinal pseudo-obstruction
- Slow transit constipation
- Chronic constipation, including constipation of aging
- Pathogen-induced diarrhea
- IBS
- Autoimmune enteric neuritis, paraneoplastic syndrome
- Enteric neuritis of unknown etiology
- Internal anal sphincter achalasia

**Neuropathies secondary to, or associated with, other disease states**

- Diabetic gastroparesis and other diabetes-related motility disorders
- Enteric neuropathy of Parkinson disease
- Enteric neuropathy of prion disease
- Enteric neuropathies associated with mental retardation or other central nervous system disorders
- Ischemic enteric neuropathy, such as in ischemic colitis

**Iatrogenic or drug-induced neuropathies**

- Disorders initiated by antineoplastic drugs (including vinca alkaloids and cisplatin)
- Postoperative ileus
- Neuropathy of ischemia, reperfusion injury, such as that associated with intestinal transplantation
- Opioid-induced constipation (usually caused when opioids are used to treat chronic pain)

**The esophagus and stomach**

*Esophageal body*

The esophageal body has one primary physiologically relevant pattern of movement—esophageal peristalsis.<sup>17</sup> The nerve circuits for motor programs of propulsive activity in the upper part of the esophagus, where the contractile elements are composed of striated muscle, are in the medulla oblongata of the CNS. These circuits relay through the nucleus ambiguus, which contains the cell bodies of the motor neurons that innervate the motor endplates of the striated muscle cells.<sup>17,18</sup> Although numerous ganglia form an ENS of conventional appearance in the striated muscle esophagus, the CNS provides the primary control of propulsion; indeed, if the vagi are severed the upper part of the esophagus is paralyzed and food is no longer propelled.<sup>19</sup> Nevertheless, processes of enteric neurons end at motor endplates in close proximity to the striated muscle cells and vagal nerve endings, and cause presynaptic inhibition of transmission from the vagal motor neurons.<sup>20,21</sup> Through the course of evolution, and with advanced encephalization in vertebrates, this part of the ENS is hypothesized to have lost an essential role that it had in more primitive animals.<sup>2</sup> Although nerve fibers that innervate the smooth muscle of the lower esophagus have their cell bodies in enteric ganglia, peristalsis in this region is also coordinated by the CNS. Indeed, the enteric ganglia of the smooth muscle esophagus are directly innervated by neurons of the dorsal motor nucleus of the vagus, and lesion of this nucleus impairs the motility patterns of the smooth muscle esophagus.<sup>17</sup>

*Lower esophageal sphincter*

A site of major control by enteric neurons is encountered at the lower esophageal sphincter (LES). Peak pressures during gastric mixing contractions exceed resting pressures in the body of the esophagus. Thus, the esophago-gastric junction, of which the LES is a major component, has the important role of limiting reflux of the corrosive contents of the stomach into the esophageal body. Failure of this guarding results in reflux esophagitis and esophageal mucosal damage, which can initiate metaplastic change (that is, Barrett esophagus).<sup>22</sup>

Two reflexes regulate the LES to enable passage of food into the stomach and to restrict reflux. First, a descending inhibitory reflex relaxes the sphincter when a bolus of food enters the last part of the esophageal body and its intraluminal pressure is raised, initiating peristaltic contractions of the esophagus to force food into the stomach. This reflex has a predominant vagovagal component and swallowing-induced reflex relaxation is inhibited by cooling the vagus nerve.<sup>23</sup> Sphincter relaxation, however, still occurs in response to distension after vagal block, indicating that a local reflex can be elicited.<sup>23</sup> As is true in many regions of the gastrointestinal tract, including the pyloric sphincter and the internal anal sphincter, the inhibitory reflex is required for the passage of solid or semisolid material and the final motor neurons of the reflex are enteric inhibitory neurons that release nitric oxide as their primary transmitter (Table 1). Of note,

**Table 1** | Multiple transmitters of neurons that control digestive function\*

Type of neuron	Primary transmitter	Secondary transmitters, modulators	Other neurochemical markers	Study
Enteric excitatory muscle motor neuron	ACh	Tachykinin, enkephalin (presynaptic inhibition)	Calretinin, $\gamma$ -aminobutyric acid	Brookes <i>et al.</i> (1991); <sup>76</sup> Holzer & Holzer Petsche (1997); <sup>77</sup> Grider (2003) <sup>78</sup>
Enteric inhibitory muscle motor neuron	Nitric oxide	VIP, ATP or ATP-like compound, carbon monoxide	PACAP, opioids	Fahrenkrug <i>et al.</i> (1978); <sup>79</sup> Costa <i>et al.</i> (1992); <sup>80</sup> Sanders & Ward (1992); <sup>81</sup> Xue <i>et al.</i> (2000) <sup>82</sup>
Ascending interneuron	ACh	Tachykinin, ATP	Calretinin, enkephalin	Brookes <i>et al.</i> (1991) <sup>83</sup>
ChAT, NOS descending interneuron	ATP, ACh	ND	Nitric oxide, VIP	Young <i>et al.</i> (1995); <sup>84</sup> Brookes (2001) <sup>85</sup>
ChAT, 5-HT descending interneuron	ACh	5-HT, ATP	ND	Furness & Costa (1982); <sup>86</sup> Monro <i>et al.</i> (2002); <sup>87</sup> Gwynne & Bornstein (2007) <sup>88</sup>
ChAT, somatostatin descending interneuron	ACh	ND	Somatostatin	Gwynne & Bornstein (2007); <sup>88</sup> Portbury <i>et al.</i> (1995) <sup>89</sup>
Intrinsic sensory neuron	ACh, CGRP, tachykinin	ND	Calbindin, calretinin, IB4 binding	Grider (2003); <sup>78</sup> Gwynne & Bornstein (2007); <sup>88</sup> Li & Furness (1998); <sup>90</sup> Johnson & Bornstein (2004) <sup>91</sup>
Interneurons supplying secretomotor neurons	ACh	ATP, 5-HT	ND	Suprenant (1984); <sup>92</sup> Monro <i>et al.</i> (2004) <sup>93</sup>
Noncholinergic secretomotor neuron	VIP	PACAP	NPY (in most species)	Cassuto <i>et al.</i> (1981); <sup>94</sup> Banks <i>et al.</i> (2005) <sup>95</sup>
Cholinergic secretomotor neuron	ACh	ND	Calretinin	Brookes <i>et al.</i> (1991); <sup>83</sup> Keast <i>et al.</i> (1985) <sup>96</sup>
Motor neuron to gastrin cells	GRP, ACh	ND	NPY	Holst <i>et al.</i> (1987); <sup>97</sup> Weiget <i>et al.</i> (1997) <sup>98</sup>
Motor neurons to parietal cells	ACh	Potentially VIP	ND	Nilsson <i>et al.</i> (1972); <sup>99</sup> Feldman <i>et al.</i> (1979) <sup>100</sup>
Sympathetic neurons, motility inhibiting	Noradrenaline	ND	NPY in some species	Finkleman (1930); <sup>101</sup> Macrae <i>et al.</i> (1986) <sup>102</sup>
Sympathetic neurons, secretion inhibiting	Noradrenaline	Somatostatin (in guinea pig)	ND	Costa & Furness (1984) <sup>103</sup>
Sympathetic neurons, vasoconstrictor	Noradrenaline, ATP	Potentially NPY	NPY	Dresel & Wallentin (1966); <sup>104</sup> Furness (1971); <sup>105</sup> Furness <i>et al.</i> (1983) <sup>106</sup>
Intestinofugal neurons to sympathetic ganglia	ACh	VIP	Opioid peptides, CCK, GRP	Crowcroft <i>et al.</i> (1971); <sup>107</sup> Dalsgaard <i>et al.</i> (1983); <sup>108</sup> Love & Szurszewski (1987) <sup>109</sup>

\*This field continues to advance as improved pharmacological and other tools are developed. Some of the information provided here will therefore undoubtedly be superseded in the future. Further details of postsynaptic receptors have been described elsewhere.<sup>2,88</sup> Abbreviations: 5-HT, 5-hydroxytryptamine; ACh, acetylcholine; CCK, cholecystokinin; ChAT, choline acetyltransferase; CGRP, calcitonin gene-related peptide; GRP, gastrin releasing peptide; ND, not determined; NPY, neuropeptide Y; NOS, nitric oxide synthase; PACAP, pituitary adenylyl-cyclase activating peptide; VIP vasoactive intestinal peptide.

although enteric neurons utilize multiple transmitters,<sup>2</sup> each class of neuron has a primary transmitter, which, in this case, is nitric oxide. Nitric oxide is formed in neurons by neuronal nitric oxide synthase. A defect in nitric oxide production results in a failure of esophageal propulsion (termed esophageal achalasia).<sup>11</sup> Moreover, failure of nitric-oxide-mediated transmission from enteric neurons in other parts of the gastrointestinal tract also causes defects in propulsion.<sup>24</sup> For example, in the stomach this defect causes gastroparesis, at the pyloric sphincter it causes hypertrophic pyloric stenosis and in the final part of the colon, deficiency of nitric oxide transmission is involved in internal anal sphincter achalasia and Chagas disease.<sup>24</sup> In addition, increased pressure in the stomach initiates a reflex constriction of the LES that limits reflux of gastric contents which, like the descending inhibitory reflex, is mediated by a vagovagal reflex pathway that passes through the brain stem.<sup>25,26</sup> Failure of the maintained closure of the LES when intra-gastric pressure exceeds intraesophageal pressure results in GERD.<sup>27</sup> In summary, although a structurally well-developed ENS exists in the esophageal wall, propulsive activity of the esophagus and the activity of the LES are

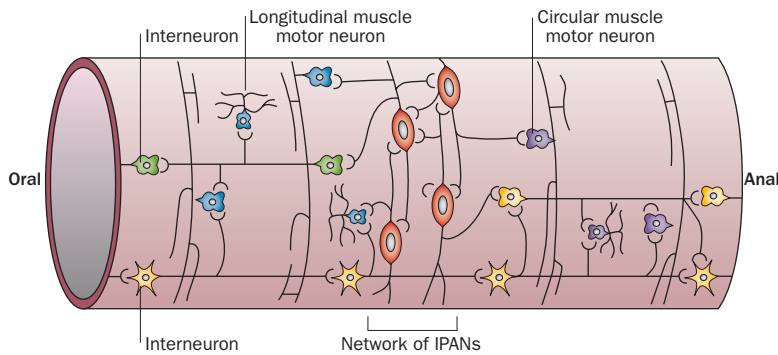
controlled through motor pattern generators in the brain stem and reflexes that depend on centers in the CNS.

#### Stomach

The stomach is similar to the esophagus in that much of its neural control is dependent on vagovagal reflexes.<sup>2</sup> However, the rhythmic contractile waves that pass caudally over the stomach to knead and propel the contents are generated in the muscle through the activity of interstitial cells of Cajal<sup>28</sup> and coordinated gastric muscle activity that is dependent on the ENS is difficult to demonstrate.<sup>2,29,30</sup> By contrast, propulsive activity in the small intestine and colon is dependent on ENS reflexes.

#### Small intestine

The ability of the small intestine to function when isolated from the CNS was demonstrated over a century ago.<sup>1</sup> The small intestine is now known to be dependent on the ENS to direct its various patterns of movement: rapid orthograde propulsion of contents (peristalsis), mixing movements (segmentation), slow orthograde propulsion (the migrating myoelectric complex, MMC) and retro-pulsion (expulsion of noxious substances associated with



**Figure 3** | Nerve circuits for control of motility in the small intestine. Nerve circuits were identified from studies in the guinea pig small intestine and similar component neurons have been identified in the small intestine of other species, including humans, and in the large intestine. Nevertheless, this figure is a simplified circuit diagram showing the major circuit features that have been identified. Networks of interconnected intrinsic sensory neurons (IPANs; red) detect mechanical distortion and luminal chemistry. These synapse with descending (yellow) and ascending (green) interneurons, excitatory muscle motor neurons (blue) and inhibitory muscle motor neurons (purple). Abbreviation: IPAN, intrinsic primary afferent neuron. Modified with permission from Furness, J. B. *The Enteric Nervous System* (Blackwell, Oxford, 2006).

vomiting).<sup>31</sup> To orchestrate these movement patterns, the state of the intestine is sensed and appropriate motor patterns are generated. The structural organization of the circuits that detect the state of the small intestine, integrate the information and direct the activities of motor neurons has now been determined (Figure 3).<sup>2</sup>

Original observations suggested that the ENS contains intrinsic sensory neurons (also referred to as intrinsic primary afferent neurons, IPANs), interneurons and both excitatory and inhibitory motor neurons that innervate the muscle.<sup>1</sup> However, identification of these neurons by their shapes, projections to targets, neurochemistries and cell physiologies has only occurred in the past decade. IPANs were first identified as large neurons with multiple, usually 2–6, long processes (type II morphology) that respond to changes in luminal chemistry, mechanical distortion of the mucosa and direct mechanical distortion of their processes in the external musculature.<sup>32–36</sup> Since that time, mechanical distortion has also been shown to excite some unipolar neurons in the enteric nerve circuits,<sup>37–39</sup> indicating that reflexes are not uniquely initiated through type II neurons. Cell bodies of multipolar IPANs are 10–30% of neurons in the submucosal and myenteric ganglia of the small and large intestines. The long axon-like processes of these neurons conduct action potentials. Moreover, action potentials that are initiated in one process traverse the cell bodies and are conducted along other processes to synaptic connections with IPANs, interneurons and motor neurons (Figure 3).<sup>36,40</sup> Consistent with the motor functions of the esophagus being controlled from or through the brain stem, type II neurons are not found in the esophagus and are rare in the stomach.<sup>2</sup>

Enteric IPANs have many features in common with small diameter primary afferent neurons of the dorsal root ganglia (DRG), which connect in the CNS. These features include containing peptides that are typical

of small-diameter afferent neurons (such as substance P and calcitonin gene-related peptide), having tetrodotoxin-resistant voltage-dependent Na<sup>+</sup> and high-voltage-activated Ca<sup>2+</sup> currents, isolectin B4 binding and unmyelinated axons.<sup>32,41</sup> As with small-diameter DRG primary afferents, the enteric IPANs are polymodal and some, at least, are nociceptors.<sup>32,41</sup> However, unlike DRG neurons, the multipolar IPANs connect with each other to form self-reinforcing networks (Figure 3).<sup>42</sup> The similarities of IPANs and DRG primary afferent neurons that supply visceral organs suggest that primary afferent neurons of neural crest origin have similar phenotypes independent of the final location of their cell bodies. Thus, these characteristics can be used to identify IPANs in other species, for example in humans<sup>3,43</sup> and large mammals<sup>44</sup> where physiological characterization is not always feasible. Structural identification of these neurons has relevance for the investigation of both human and animal enteric neuropathies.

Although the neural circuits are now characterized, the mechanisms within the integrative circuitry through which one pattern of activity is converted to another are not yet defined. Nevertheless, signals that trigger changes in patterns of movement have been identified. For example, fatty acids added to the luminal surface convert propulsive contractile activity to mixing movements, through a neural mechanism.<sup>45</sup> Conversion from one pattern to another can also be achieved with drugs that target enteric neurons. For example, TRAM34-mediated inhibition of the intermediate conductance, calcium-activated, potassium IKCa channel, which underlies the slow hyperpolarization that follows the action potential in IPANs, changes propulsive movements to mixing movements, possibly by altering the timing of action potential firing in IPANs.<sup>46,47</sup>

The MMC traverses the small intestine periodically between meals, in humans approximately every 90 min.<sup>48</sup> Its passage is slow, ~1–4 cm per minute in humans, and its suggested role is to remove residual contents from the intestine after a meal has been digested (and thus it has been called the intestinal housekeeper), thereby reducing bacterial overgrowth in the small intestine.<sup>31</sup> The MMC is dependent on the ENS for its progress along the intestine, as shown by its inhibition by the local infusion of tetrodotoxin<sup>49</sup> or by the nicotinic blocker, hexamethonium.<sup>50</sup>

The detection of noxious agents (nociception) is important to initiate reactions that expel pathogens and potentially injurious chemicals. Intrinsic retropropulsive reflexes in the small intestine force the contents back to the stomach in association with vomiting,<sup>51,52</sup> and in the large intestine pathogens activate the ENS to cause large propulsive contractions and copious fluid secretion.<sup>53,54</sup> Under non-pathological conditions fluid secretion is also coupled with contractions of the circular muscle of the bowel wall.<sup>55</sup>

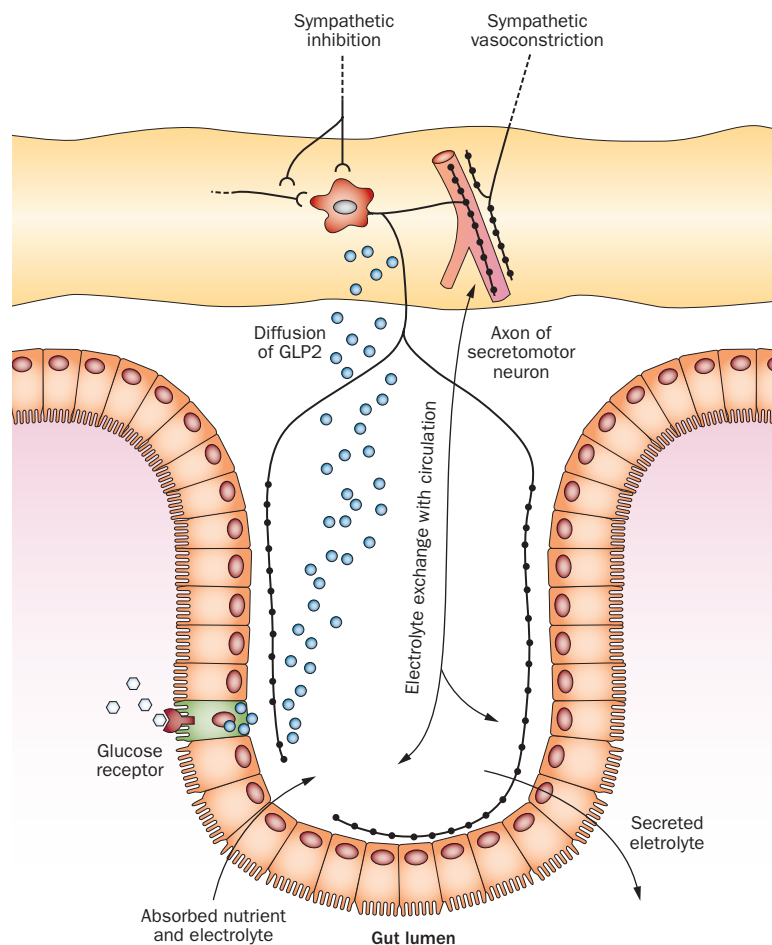
### Large intestine

Colonic motility is dependent on the ENS for propulsion of contents. Indeed, colonic motility fails when the ENS of the distal colon and rectum is congenitally absent in

patients with Hirschsprung disease,<sup>7</sup> when it degenerates in later life, as in Chagas disease<sup>8</sup> or is compromised in other forms of enteric neuropathies. Nevertheless, the CNS directs ENS activity in the colorectum. In healthy individuals, the propulsive reflexes of the distal colon and rectum are regulated by central control centers that maintain fecal continence and, when it is appropriate, trigger defecation through central commands that are relayed through the defecation center in the lumbosacral spinal cord (Figure 1).<sup>16,56</sup> Indeed, direct stimulation of the defecation center in the spinal cord causes coordinated emptying of the colon, via the ENS.<sup>57</sup> Voluntary controls of defecation (both inhibition and facilitation) are lost if cortico–spinal connections to the defecation centers are severed by spinal injury.<sup>58</sup> Nevertheless, if the defecation center remains intact after spinal injury it can be stimulated to command the ENS pathways for bowel emptying.<sup>59</sup> Indeed, a ghrelin receptor agonist is now being developed for use in clinical practice in patients with spinal injuries.

### Control of fluid movement

Movement of fluid between the lumen of the intestine and the body fluid compartments is tightly regulated. More than two blood volumes cross the mucosal epithelial surfaces each day, and disruption of fluid transport regulation, such as that which occurs during cholera intoxication, is life-threatening. One reason for the large flux is that the absorption of sugars (monosaccharides) and amino acids is through cation-coupled transporters. Thus, when glucose is absorbed through the sodium–glucose linked transporter it is internalized with a sodium ion and counter ions, the majority of which are chloride ions. Absorption of 100 g glucose is estimated to equate to absorption of 1.8 liter of water.<sup>2,60</sup> Enteric reflexes, through activation of secretomotor neurons, move water and electrolytes from the interstitium of the lamina propria to the lumen (Figure 4). The water and electrolytes are drawn from both the circulation and the absorbed fluid. Enteric secretomotor reflexes cannot act in isolation and are modulated to control whole body fluid balance. This control is exerted through blood volume and blood pressure detectors that change the activity of two sympathetic pathways, vasoconstrictor pathways and secretomotor inhibitory pathways (Figure 4).<sup>2,61</sup> The sweet taste receptor, which is the enteric receptor for glucose, is composed of T1R2 and T1R3 subunits and located on enteroendocrine L cells.<sup>62,63</sup> Stimulation of these cells releases a group of endocrine hormones: glucagon-like peptide (GLP)1 and GLP2, oxyntomodulin and peptide YY (PYY). Receptors for GLP2 are on non-cholinergic secretomotor neurons, which are activated by this hormone.<sup>63,64</sup> Thus, activation of the enteric receptor for glucose by glucose or artificial sweeteners stimulates secretomotor neurons to return water and electrolytes to the lumen (Figure 4). The other endocrine products of L cells are involved in a range of functions, including modulation of fluid secretion, inhibition of gastric emptying, increasing satiety, stimulation of epithelial cell growth and regulation of insulin secretion.<sup>64–66</sup>



**Figure 4** | Neural control of transmucosal water and electrolyte movement in the small intestine. Neural circuitry has an essential role in balancing local fluid fluxes and in whole body water and electrolyte balance. Large volumes of fluid are absorbed from the lumen with nutrients, such as glucose, which are transported by ion-coupled transporters. These fluids are returned through secretomotor reflexes. Luminal glucose stimulates secretomotor neurons via the glucose receptor on L-type enteroendocrine cells (green), which release the hormone GLP2, which diffuses through the lamina propria to activate secretomotor neurons. The balance of this fluid exchange is modulated by sympathetic vasoconstrictor and secretomotor inhibitory pathways. Activity in these sympathetic pathways, which inhibit secretion and reduce local blood flow, is determined by whole body fluid status, which includes sensory detection through volume detectors, baroreceptors and osmoreceptors. Abbreviation: GLP2, glucagon-like peptide 2. Modified with permission from Furness, J. B. *The Enteric Nervous System* (Blackwell, Oxford, 2006).

The fine control of fluid balance through local (ENS) and systemic (sympathetic) reflexes is thrown into chaos when the lumen contains an excess of certain pathogens or their toxins, including cholera toxin, rotavirus and pathogenic *Escherichia coli*, which activate enteric secretomotor neurons.<sup>12,67</sup> In mild cases, diarrhea is induced that helps to expel the pathogens and their toxic products. However, high levels of pathogens or toxins overwhelm the intestine and pathological, life-threatening diarrhea can develop.

### Chemistry of neurotransmission

The enteric neurons and the extrinsic neurons that connect to the gastrointestinal tract release multiple transmitters (Table 1). In the majority of cases a primary

transmitter and subsidiary transmitters or transmission modulators have been identified. The synthesis pathways, release mechanisms, transmitter metabolic pathways and the receptors for the transmitters are obvious targets for drug therapy, as well as for studies of the physiological relevance of the transmitter systems. For example, cholinergic transmission has been targeted for slowed gastrointestinal transport<sup>68</sup> and alvilopam, the peripherally acting opioid receptor antagonist, has been used for treating post-operative ileus.<sup>69</sup> Other receptors that have been targeted are those for tachykinins (NK1 and NK3 receptors) in IBS and 5-HT (5-HT<sub>2B</sub>, 3 and 4 receptors) in constipation and IBS,<sup>70–72</sup> but therapeutic advances based on the burgeoning knowledge of the ENS and its neurochemistry have been less promising than might be expected.<sup>71,73</sup>

### Enteroendocrinology

Many functions of the digestive system, and functions related to digestion, such as satiety, involve both enteric innervation and the endocrine system of the digestive tract (called the gastroenteropancreatic endocrine system). In fact, most aspects of gastrointestinal control involve both neurons and endocrine cells. Similarly to the ENS, the gastroenteropancreatic system is extensive, consisting of approximately 30 endocrine cell types that secrete ~100 messenger molecules.<sup>74</sup> Although a discussion of all aspects of the gastroenteropancreatic system is beyond the scope of this Review, of note, the mucosal endings of sensory neurons are separated from the luminal contents by a continuous epithelial lining. Thus, luminal contents are detected by receptors on enteroendocrine cells, which release messenger molecules from their basolateral surfaces to activate enteric, vagal and spinal sensory neurons.<sup>34,63,75</sup> A good example of neural and endocrine integration is the control of gastric acid secretion. The primary endocrine hormones involved in control of acid secretion are gastrin released from antral G-type enteroendocrine cells, histamine released from

enterochromaffin-like cells and somatostatin released from gastric mucosal D cells.<sup>15</sup> However, the parietal cells are influenced both directly and indirectly by cholinergic neurons with cell bodies in the gastric ENS and gastrin release is controlled by gastrin-releasing peptide—a transmitter released from enteric neurons that innervate G cells.<sup>2,15</sup>

### Conclusions

The ENS is one component of the neural control system of the digestive tract. It works in concert with the CNS, integrative pathways that pass through sympathetic ganglia and the gastroenteropancreatic endocrine system. In the small and large intestines, the ENS contains full reflex pathways that are essential to direct the movements of these parts of the digestive tract and to control fluid movement between the gut lumen and body compartments. The control of fluid transport represents a case of close integration between the CNS (via sympathetic pathways) and the ENS. The primary control centers for the smooth muscle esophagus, LES and stomach are in the CNS, but the connections to effectors in these organs is through the ENS. The striated muscle esophagus is controlled through circuits in the medulla oblongata and the muscle is directly innervated by vagal neurons with cell bodies in the CNS. Thus, neurogastroenterology is the science of integrated ENS and CNS control. The range of enteric transmitter systems and their receptors provide the basis for numerous future therapeutic targets.

#### Review criteria

Full-text articles were selected from the author's personal library collected over the past 40 years and from searches of the PubMed database. The following search terms were used: "enteric nervous system", "enteric neurons", "gastroenterology", "motility", "intestinal secretion", "stomach" and "small intestine".

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