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## Gut feelings: the emerging biology of gut–brain communication

**Emeran A. Mayer**

Center for Neurobiology of Stress, Division of Digestive Diseases, Departments of Medicine, Physiology and Psychiatry, David Geffen School of Medicine at University of California, Los Angeles, CHS 47-122 10833 Le Conte Avenue, Los Angeles, California 90095-7378, USA

Emeran A. Mayer: emayer@ucla.edu

### Abstract

The concept that the gut and the brain are closely connected, and that this interaction plays an important part not only in gastrointestinal function but also in certain feeling states and in intuitive decision making, is deeply rooted in our language. Recent neurobiological insights into this gut–brain crosstalk have revealed a complex, bidirectional communication system that not only ensures the proper maintenance of gastrointestinal homeostasis and digestion but is likely to have multiple effects on affect, motivation and higher cognitive functions, including intuitive decision making. Moreover, disturbances of this system have been implicated in a wide range of disorders, including functional and inflammatory gastrointestinal disorders, obesity and eating disorders.

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A major scientific breakthrough in understanding the interaction of the nervous system with the digestive system occurred with the discovery of the so-called enteric nervous system (ENS) in the middle of the nineteenth century<sup>1–4</sup> (BOX 1). Even though it is now considered the third branch of the autonomic nervous system, the ENS has been referred to as the ‘second brain’, based on its size, complexity and similarity — in neurotransmitters and signalling molecules — with the brain<sup>5</sup>. Even before the discovery of the ENS, the importance of interactions between the brain and the digestive system in health and disease has been recognized for many centuries, and has been studied by prominent psychologists, psychiatrists and physiologists during the later part of the nineteenth and earlier part of the twentieth centuries (reviewed in REF. 6). Both top-down modulation of gastrointestinal function by stress and emotions<sup>7</sup>, and bottom-up signalling from visceral afferents to the brain in abdominal pain syndromes, as well as possible emotion regulation<sup>8</sup> have been reported by early investigators (BOX 2). This topic has received increased attention during the past two decades, largely owing to a series of independent but converging scientific discoveries from various fields of research, including enteric neuroscience (reviewed in REF. 1), neuroimaging (reviewed in REF. 9), intestinal microbiology and host microbial interactions (reviewed in REFS 10,11), and most recently, microbial gut–brain signalling (reviewed in REFS 12–14). This Review discusses the emerging role of integrated bidirectional signalling between the brain and the gut in homeostasis and disease, as well as possible consequences for higher-level executive functions and emotional states, with a primary focus on gut to brain signalling.

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**Further Information:** Emeran A. Mayer's homepage: <http://www.uclacns.org>

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## Brain to gut signalling during homeostasis

The brain communicates to the viscera, including the gastrointestinal tract, through multiple parallel pathways, including the two branches of the autonomic nervous system (ANS), the hypothalamic–pituitary–adrenal (HPA) axis and the sympatho–adrenal axis (modulating the gut-associated lymphoid tissue), and descending monoaminergic pathways (modulating gain of spinal reflexes and dorsal horn excitability). Two key subcortical structures that generate these outputs are the hypothalamus and the amygdala. They receive inputs from a network of cortical regions (the medial prefrontal cortex (PFC) network<sup>15,16</sup>), including subregions of the medial PFC and anterior cingulate cortex (ACC). Whereas ventral ACC regions (for example, subgenual cingulate cortex and Brodmann area 25 (BA25)) project primarily to the medullary vagal complex, pregenual ACC regions (for example, BA24) project to the periaqueductal grey (PAG)<sup>17</sup>. The medial PFC network receives input from a lateral PFC and orbitofrontal cortex (OFC) network that provides integrated multisensory information about the representation of complex homeostatic body states, including those related to gut homeostasis, food intake and visceral pain. Outputs from subregions of the medial network, amygdala and hypothalamus are integrated into distinct motor patterns within the mesencephalic PAG<sup>18</sup>. The most caudal component of the central autonomic networks is represented by pontine and medullary nuclei, including the serotonergic raphe nuclei, the locus coeruleus complex (including Barrington's nucleus) and the dorsal vagal complex. This system of parallel outflows from cortico–limbic–pontine networks, which is engaged by distinct homeostatic states, has been referred to as the emotional motor system (EMS) and consists of integrated motor autonomic, neuroendocrine and pain modulatory components<sup>19,20</sup>. The medial component of the EMS (including the midline raphe nuclei) is thought to have a role primarily in the tonic modulation through serotonergic, noradrenergic and opioidergic descending spinal pathways of the gain of various spinal reflexes involved in the regulation of gastrointestinal functions, as well as in the modulation of dorsal horn neuronal excitability, to regulate pain sensitivity. An example of the gut-related engagement of the medial component of the EMS is the observation that food consumption inhibits pain-related behaviours in the rat, an analgesia mechanism related to the engagement of descending serotonergic pain modulation pathways<sup>21</sup>. Descending opioid-dependent pain modulation pathways are also engaged by other behavioural and motivational states, including vigilance and fear<sup>22</sup>. By contrast, the lateral system (including subregions of the PAG, the central nucleus of the amygdala, bed nucleus of the striae terminalis, lateral hypothalamus and pontine locus coeruleus complex) may play a part in executing distinct regional motor patterns of the viscera mediated by the engagement of function-specific subsets of sympathetic and parasympathetic pathways.

ANS output can be triggered reflexively by ascending interoceptive signals from the gut, by descending cognitive or emotional influences or in response to external or internal demands (BOX 3). For example, top-down modulation can override local (for example, ENS-based) reflex function in the context of threats to body homeostasis (haemorrhagic shock), severe environmental stressors<sup>23</sup> or during strong emotions such as fear, anger and sadness<sup>24,25</sup>.

Outputs from medullary pontine nuclei project to function-specific sympathetic preganglionics located in the intermediolateral column of the spinal cord, and projections to the medullary vagal complex and the pontine Barrington's nucleus modulate function-specific vagal and sacral parasympathetic outflows, respectively<sup>26</sup>. Preganglionic parasympathetic neurons are viscerotopically organized in the dorsal motor nucleus of the vagus and are modulated by a variety of hormonal and humoral substances<sup>27,28</sup>.

## Effects on the gut

The sympathetic innervation of the gastrointestinal tract and its role in the modulation of gastrointestinal function has been extensively reviewed<sup>1,28,29</sup>. It has been divided into subclasses of postganglionic vasoconstrictor neurons, secretion inhibiting neurons and motility inhibiting neurons. The overall effect of sympathetic outflow to the gut is inhibitory, slowing gastrointestinal transit and secretion. This inhibitory effect is largely accomplished by inhibitory modulation of cholinergic transmission and by a stimulatory effect on smooth muscle in sphincteric regions. Another subset of sympathetic postganglionics is involved in mucosal immune modulation<sup>30</sup> and possibly in the modulation of interactions between the micro-flora and the mucosa (reviewed in REFS 12,31). Although the best experimental evidence for such communication between the sympathetic nervous system, lymphocytes and macrophages comes from the spleen<sup>32</sup>, evidence for sympathetically mediated immune modulation in the gut has been shown both in Peyer's patches and in the non-follicular mucosa in close proximity to different classes of immune cells, including dendritic cells, mast cells and B lymphocytes (reviewed in REF. 31). In addition, evidence of a noradrenaline-mediated reduction in the expression of toll-like receptors (TLRs) by intestinal epithelial cells has been reported<sup>33</sup>. Finally, noradrenaline-mediated modulation of microbial virulence has been reported for a number of pathogens<sup>31</sup>, even though the mechanisms by which intraluminal bacteria are exposed to noradrenaline that is released from sympathetic nerve terminals is unclear.

The parasympathetic innervation of the gastrointestinal tract has been studied intensively (for reviews see REFS 28,34). It is comprised of the vagal and sacral parasympathetic divisions, which innervate foregut and hindgut structures, respectively. Function-specific vagal motor neurons provide input to the stomach, small intestine and proximal portion of the colon. Excitatory vagal input occurs to ganglia within the ENS to mediate vago-vagal motor reflexes and the cephalic phase of gastric acid secretion, to gastrin- and somatostatin-containing enteroendocrine cells and histamine releasing enterochromaffin cells, and to enterochromaffin cells to mediate 5-hydroxytryptamine (5-HT) release<sup>35</sup>. While vago-vagal reflexes are the primary neural mechanism to regulate gastric function, such extra intestinal reflexes play a lesser part (compared to intrinsic reflexes) in the modulation of intestinal function. Vagal modulation of macrophage activation through nicotinic acetylcholine receptors has been reported as part of a vago-vagal anti-inflammatory reflex<sup>36</sup>.

Top-down engagement of subsets of function-specific postganglionic sympathetic and parasympathetic neurons are likely to mediate reported emotion-related patterns of regional changes in motor, secretory and possibly immune activity in the gastrointestinal tract<sup>37,38</sup>, which may be viewed as analogous to distinct emotion-related facial expressions and body postures (mediated by the somatic branch of the EMS). These emotion-related changes in peripheral target cells will directly or indirectly (for example, through changes in smooth muscle activity) influence the interoceptive feedback to the brain, possibly contributing to the characteristic prolonged duration of many emotional states, which can outlast the initiating event by hours. Prolonged alterations in ANS output to the gut is likely to induce changes in peripheral target cells — such as downregulation of adrenergic receptors on immune cells<sup>39</sup>, reduced expression of TLRs<sup>33</sup> on epithelial cells or changes in primary afferent neurons<sup>40</sup> — changing the gain of gut to brain signalling chronically and possibly resulting in the remodelling of brain regions that receive this enhanced input<sup>41</sup>. Such lasting changes in brain-gut signalling may be associated with tonic ANS dysfunction, which is in turn associated with altered emotional states such as anxiety disorders or depression.

## Gut to brain signalling during homeostasis

There are between 200 and 600 million neurons in the human ENS, which is equal to the number of neurons in the spinal cord<sup>1</sup>. They have been classified on the basis of their morphology, electrophysiological properties and chemical coding into distinct classes of functional specific neurons, including several classes of afferents<sup>1,42</sup>. The size and complexity of the ENS is not surprising when considering the challenges posed by the interface of the organism with its luminal environment: it interfaces closely with our largest body surface (the intestinal surface area, which is approximately 100 times larger than the surface area of the skin), with the largest population of commensal microorganisms of all body surfaces (100 trillion microorganisms from 40,000 species with 100 times the number of genes in the human genome<sup>43</sup>), with the gut-associated immune system (containing two-thirds of the body's immune cells) and with thousands of enteroendocrine cells (containing more than 20 identified hormones). These unparalleled relationships between the gastrointestinal tract and the brain, with multiple bidirectional and often interacting interoceptive communication systems, emphasize the importance of this system in the maintenance of homeostasis. There are three basic mechanisms by which sensory information is encoded in the gut: by primary afferent neurons, by immune cells and by enteroendocrine cells (FIG. 1a).

### Neuronal signaling

Based on the location of their cell body, the afferent neurons that innervate the gut are divided into extrinsic (spinal and vagal afferents) as well as several classes of intrinsic, primary afferents (IPANs)<sup>44</sup> (FIG. 1a). IPANs greatly outnumber the extrinsic connections, with an estimated 100 million IPANs versus 50,000 extrinsic afferent nerve processes in the human gut<sup>1</sup>. Both intrinsic and extrinsic primary afferents show mechano- and chemosensitivity to both physiological and noxious mechanical stimuli<sup>45</sup>. Luminal signals do not influence afferent nerve terminals directly, but act through intermediate cells in the lamina propria (including enteroendocrine cells)<sup>46</sup>. Both extrinsic and intrinsic primary afferents provide input to multiple reflex loops to optimize gut function and maintain gastrointestinal homeostasis during internal perturbations (BOX 3). Subpopulations of modality-specific extrinsic primary afferents synapse with dorsal horn neurons in lamina I of the spinal dorsal horn or with cell bodies in the vagal nucleus tractus solitarius (NTS), providing the brain with a wide range of gut-related information. There are differences between the stomach and the intestine in the relative importance of intrinsic and extrinsic reflex regulation: whereas gastric function is predominantly regulated by vago-vagal reflexes, which are modulated within the dorsal vagal complex by cortical, humoral and endocrine signals, intestinal function relies primarily on intrinsic reflex regulation, mediated by IPANs and enteric motoneurons<sup>1</sup>.

IPANs are connected with each other through excitatory synapses, forming extensive self-reinforcing networks<sup>47</sup>. Such networks are likely to amplify local signals that arise through direct activation of IPANs by an afferent stimulus. Some IPANs are polymodal, for example, responding to both chemical stimuli arising from entero-chromaffin cells as well as to mechanical stimuli that result from gut contractions and distensions. Whereas IPANs feed this information into a variety of local intramural reflexes that regulate and integrate gut functions such as propulsion, secretion and blood flow (but do not communicate directly with the CNS), extrinsic afferents provide afferent information to mesenteric, spinal and supraspinal reflex loops involving larger distances of the bowel and different regions of the gastrointestinal tract.

The fibres of the mechanosensitive extrinsic afferents that innervate the gastrointestinal tract end within the mesentery and longitudinal and circular muscle sheets, within the enteric

ganglia<sup>48–50</sup>, or in close proximity to interstitial cells of Cajal<sup>51</sup>. These afferent terminals are sensitive to distension, contractions or movement of the gut through activation of a variety of mechano-sensitive ion channels, including members of the transient receptor potential (TRP) family (TRPV1, TRPV2, TRPV3 and TRPV4, and TRPA1) and acid-sensing channels (ASICs)<sup>52</sup>. Subclasses of extrinsic afferents have been identified based on their mechanical and electro-physiological response properties, including low- and high-threshold mechanosensitive vagal and spinal afferents<sup>53–55</sup>. A large number of such afferents do not respond to mechanical stimuli under normal conditions but develop mechanosensitivity following inflammation<sup>56,57</sup>.

Different subpopulations of primarily vagal afferents terminate in close proximity (but without specialized synaptic connections) to enteroendocrine cells<sup>58</sup>. Their terminals are chemosensitive, responding to neuropeptides released from enteroendocrine cells in response to luminal chemical and mechanical events (FIG. 1). Different classes of vagal afferents express receptors for both orexigenic (hunger generating) as well as anorexigenic (satiety generating) gut peptides contained in specialized enteroendocrine cells distributed throughout the gastrointestinal tract, from the oral cavity to the colon. Recent evidence suggests that the expression of these receptors on vagal afferents can be altered by the nutritional state of the animal and possibly by diet<sup>59</sup>, and that the cholecystokinin 1 receptor (CCK1R; also known as CCKAR) has an important role in this process<sup>60,61</sup>. The expression of receptors for orexigenic peptides was found to be increased by fasting (increasing the signal to eat), whereas that for anorexigenic peptides was decreased. These findings demonstrate the plasticity of the vagal afferent phenotype depending on the homeostatic state of the animal, analogous to the inflammation-induced phenotype switching of mechanosensitive afferents discussed above. Terminals of some vagal afferents end in close proximity to mucosal immune cells<sup>62</sup> and have been shown to respond to a variety of signalling molecules that are released by mast cells and lymphocytes, including tryptase, histamine, 5-HT and various cytokines<sup>46</sup>.

### Endocrine and paracrine signaling

Although enteroendocrine cells make up less than 1% of all gut epithelial cells, they constitute the largest endocrine organ of the body. Based on their content, over 20 different types of enteroendocrine cells have been described<sup>63,64</sup>. Enteroendocrine cells provide the first level of integration for all chemical and certain mechanical stimuli that are related to intraluminal gut contents. Their output is involved both in the regulation of digestive functions through ENS circuits and in the regulation of CNS processes through endocrine and paracrine signalling to vagal afferents. The relative importance of paracrine (for example, vagal) versus endocrine signalling to the hypothalamus and other brain regions in the regulation of food intake remains to be determined. Some enteroendocrine cells (so-called open cells) respond to luminal contents through multiple receptors (FIG. 1b). Mechanisms of enteroendocrine and enterochromaffin cell activation include depolarization by activation of mechanosensitive cation channels located on microvilli in response to the shearing forces of moving intestinal contents<sup>64</sup>, activation by fatty acids through an acyl chain length-specific Ca<sup>2+</sup>-dependent mechanism<sup>65</sup>, presumably a G protein-coupled receptor (GPCR)<sup>66</sup>, activation by vagal efferents<sup>35</sup> as well as depolarization through G protein-coupled taste receptors<sup>67</sup>. In addition, enteroendocrine cells respond to the presence and activity of intraluminal microbial organisms through pattern recognition receptors (including TLRs), (reviewed in REFS 12,68). One example for this multimodal sensitivity is illustrated by the observation of increased release of cholecystokinin from enteroendocrine cells in response to intestinal pathogens<sup>69,70</sup>. There has been a recent surge of interest in the sensors that detect sweet and bitter sensations and amino acids in the gut, which were originally described as taste receptors in the tongue<sup>67</sup>. The taste receptor type 1 (T1R)

family comprises three members that heterodimerize to form sweet taste receptors (T1R2 and T1R3) and amino acid receptors (T1R1 and T1R3), and the T2R family includes GPCRs that are involved in bitter taste reception. As in the tongue, taste receptor signalling involves a specific G $\alpha$  subunit,  $\alpha$ -gustducin, a taste-specific signalling protein with a prominent role in bitter taste<sup>71</sup>. Each of the channels described above triggers different signal transduction mechanisms, resulting in intracellular calcium increases and peptide release, or TLR-mediated activation of NF- $\kappa$ B, activator protein 1 (AP1; also known as transcription factor AP1) and interferon regulatory factors (IRFs); which leads to cytokine release<sup>12</sup>. Crosstalk among these signalling channels might enable signals from pathogenic and commensal microorganisms to alter the response of certain enteroendocrine cells to mechanical and/or chemical stimulation<sup>59</sup>.

A particularly well-studied type of enteroendocrine cell is the 5-HT-containing enterochromaffin cell of the small and large intestine<sup>64</sup>. 95% of the organism's 5-HT is located in the gut, primarily in enterochromaffins. It is released on the basolateral side of these cells in response to shearing forces that are produced by moving luminal contents<sup>64</sup> that activate mechanosensitive cation channels located on microvilli on the apical side of enterochromaffin cells. Although mechanically released 5-HT is essential for triggering normal peristalsis and secretomotor reflexes, it is unknown if signalling to the CNS occurs during such physiological sensing and if tonic, vagally mediated 5-HT signalling to the brain is involved in modulating affective brain circuits.

### Immune-related signaling

70–80% of the body's immune cells are contained within the gut-associated lymphoid tissue, reflecting the unique challenge for this part of the immune system to maintain a homeostatic balance between tolerance and immunity in the intestine. A single layer of columnar intestinal epithelial cells (IECs) forms the barrier between the host and one of the largest and most complex microbial habitats. The fact that mucosal immune cells remain immunologically hyporesponsive to commensal bacteria but also maintain their responsiveness to pathogenic organisms, suggests that under physiological conditions, the intestinal immune system recognizes commensal bacteria and elicits basal or tonic signals in the absence of full activation of the innate and adaptive immune response<sup>72</sup>. The physical constraints that the undisturbed epithelial layer imparts on sampling the luminal space are overcome by specialized lymphoid structures (including small intestinal Peyer's patches) that are embedded in the lamina propria throughout the intestine<sup>72</sup>. Two epithelial transduction mechanisms have been identified: specialized microfold cells that can sample antigens and microorganisms, and deliver them through transcytosis to antigen processing cells within Peyer's patches, and lamina propria dendritic cells, which extend their dendrites between epithelial tight junctions, to sample the luminal environment. Similarly to enteroendocrine cells, both EICs and dendritic cells express a wide range of pattern recognition receptors, including TLRs<sup>73</sup>, emphasizing the central role of the gut in interactions with microorganisms and in discriminating between pathogens and commensals. Subsets of vagal afferent nerve terminals are in close proximity to mucosal immune cells and contain receptors for signalling molecules released from these cells, including mast cell products (proteases, histamine, serotonin and corticotropin releasing factor (CRF))<sup>62</sup> and cytokines that are released from macrophages. Immune cell products can indirectly influence the functional properties of enteroendocrine cells, as illustrated by the increased cholecystokinin secretion in a mouse model of gut inflammation owing to the release of interleukin-4 (IL-4) and IL-13 from CD4<sup>+</sup> T cells<sup>70</sup>. Similarly, changes in the properties of the enterochromaffin cell-based 5-HT signalling system in the context of gut inflammation have been described<sup>74</sup>.

## The effect on the brain

Despite the continuous stream of interoceptive signals from the gut to the brain, only a small fraction of this information is consciously perceived, usually that which requires a conscious behavioural response (for example, ingestive behaviours and defecation). However, recent evidence suggests that various forms of subliminal interoceptive inputs from the gut, including those generated by intestinal microbes, may influence memory formation, emotional arousal and affective behaviours<sup>75</sup>.

Primary vagal and spinal afferents from the gastrointestinal tract project to the NTS and the lamina I of the dorsal horn, respectively (FIG. 2). In addition, laminae V, VII and X receive additional nociceptive spinal inputs. In mammals, activity of vagal projections from the NTS and from lamina I are integrated in the parabrachial nucleus, which then provides an integrated signal to behavioural control regions of the forebrain, including the hypothalamus and amygdala<sup>76</sup>. Collaterals from these ascending vagal and spinal projections to medullary serotonergic (raphe) and catecholaminergic nuclei (including the locus coeruleus complex) provide input to ascending monoaminergic projections that can modulate cortical arousal. In non-human and human primates, there is a direct projection from lamina I and from the NTS to ventromedial thalamic nuclei. Parallel projections of lamina I afferents provide a direct thalamo–cortical pathway that activates the ACC. Based on electrophysiological and neuroanatomical data obtained in animals, Craig proposed the concept of a lamina I spino–thalamo–cortical pathway, visible in primates and well-developed only in humans, that provides a direct cortical image of the homeostatic state of the body (in the posterior and mid insular cortex) that is modulated by input from emotional and cognitive brain circuits in the ventral and dorsal portions of the anterior insula (aINS)<sup>77–80</sup> (BOX 2). Even though a close reciprocal relationship between interoceptive information (including different aspects of gut-related information) and emotion has previously been proposed<sup>8,81</sup>, the human INS — and its related brain networks (including the ACC, OFC and amygdala) — has emerged as the most plausible brain region supporting this integration. Several recent studies, including two quantitative meta-analyses<sup>82,83</sup>, have largely defined the general functional subregions of the INS, with substantial homology to the functional neuroanatomy of non-human primates. In one of these meta-analyses<sup>82</sup>, evidence for both differentiation into function-specific subregions as well as integration of multiple functions within a single subregion has been demonstrated. For example, visceral and sensory input were represented in the mid posterior aspects of the INS, and olfactory and gustatory stimuli were represented in the aINS and mid INS respectively, suggesting that distinct but closely adjacent INS subregions are responsible for the primary representation of different aspects of ingestion-related interoceptive inputs. A conjunction analysis across all of the domains revealed that the majority of all tested functions overlapped in the dorsal aINS (with the exception of basic somatosensory processes), suggesting that this brain region may be concerned with integration of food-related (olfaction and taste), interoceptive, emotional and cognitive functions.

Earlier intra-operative electrical stimulation studies<sup>84</sup> had shown that stimulation of the aINS was associated with gustatory and olfactory sensations, whereas stimulation of more posterior subregions resulted in interoceptive and somatosensory sensations. The fact that changes in gastric motility were also observed in these experiments is consistent with the homeostatic reflex concept discussed in BOX 3. In addition to the mapping of different nutrient-related functions onto the aINS, activation of a dorsal aINS subregion was seen in correlation with cognitive functions<sup>79,80</sup>, and activation in a ventral aINS subregion was seen in correlation with social–emotional functions, consistent with homologous findings in non-human primates<sup>85</sup>. A recent connectivity study based on spatiotemporal oscillations in the resting brain confirmed the existence of a ventral aINS-centered network<sup>86</sup>. Based on the close structural connectivity of different INS subregions in non-human primates<sup>85</sup>, it can be

assumed that the close spatial association of function-specific INS subregions found in human brain imaging studies allows for the necessary integration of parallel inputs related to the homeostatic state of ingestion-related functions with cognitive and emotional influences. In the case of signals related to the olfactory<sup>87</sup>, gustatory<sup>88</sup> and viscerosensory dimensions of food intake, this integration is accomplished through close connections of the aINS both to a ventral circuit (including the amygdala, ACC and OFC, and the ventral striatum) and a dorsal circuit involving the dorsolateral PFC and the dorsal striatum<sup>89</sup>.

## Emotional and executive function

### Effects of gut to brain signalling on emotional function

Although it is intriguing to speculate about the effects of enteroendocrine and enterochromaffin cell–vagal signalling to the brain as an important mechanism of emotion modulation in health and disease, the experimental literature mostly supports the hedonic effects of food intake (taste and satiation) and the aversive effects of acute mucosal inflammation or irritation by luminal toxins and pathogens. Ingesting foods that are beneficial for the organism is associated with a feeling of pleasure, whereas ingestion of potentially harmful foods elicits negative emotions, such as nausea. Gut-based neuroendocrine signalling systems (involving signalling molecules located in different classes of enteroendocrine cells) can signal satiety (including cholecystokinin, peptide YY, glucagon-like peptide 1 (GLP1)), hunger (ghrelin) and nausea (5-HT), whereas those located in mucosal immune cells (including CRF, 5-HT, proteases and cytokines) can signal pain, discomfort, nausea and fatigue. Little is known about the possible role of enterochromaffin cell–vagal signalling or the role of intestinal taste receptor activation in background emotions. However, a small number of studies in animal models show a possible role of the gut microbiota in emotional behaviour in rodents<sup>13,14,90–92</sup>, even though the mediating signalling mechanisms are poorly understood. Intraluminal infusion of nutrients has been shown to modulate flavour preferences and acceptance in rodents<sup>93</sup>. Intra-gastric infusion of fatty acids in human subjects resulted in a reduced brain response to experimentally induced sad emotions<sup>94</sup>.

### Meta-representation of gut-related feeling states

A growing literature on the topic of different sensory modalities supports the concept that actual viscerosensory (and somatosensory) stimulation, as well as expected or imagined sensory experiences, engage different regions of the INS. For example, although actual distension of the gut in healthy subjects is associated with activation of the mid INS and aINS, consistent with the reported findings about insular viscerosensory representation<sup>82</sup>, the expectation of visceral pain has been shown to be associated with activation of the aINS only<sup>95</sup>, consistent with the role of the aINS in the processing of interoceptive memories. Similar findings had previously been reported for expected somatic pain<sup>96</sup>.

Activations of aINS (and ACC) have also been reported when subjects experienced disgust, or viewed images of others experiencing disgust<sup>97</sup>, in the absence of a vagally mediated signal from the gastrointestinal tract signalling nausea. This suggests that homeostatic reflexes at the level of the aINS and ACC can be triggered in the absence of interoceptive input, presumably based on the recall of interoceptive memories while viewing an interoceptive feeling expressed by another individual or by expecting or imaging a particular feeling state.

The mechanisms and functional neuroanatomy underlying the formation of interoceptive memories of complex gut-related homeostatic states with affective and motivational dimensions (such as pain, nausea, hunger and satiation) are incompletely understood but are likely to include the encoding of such memories during development, possibly beginning



with the earliest experience of pain (hunger and abdominal discomfort) and pleasure related to food intake of the newborn. Gut to brain signalling in the newborn underlies early undifferentiated emotions and stereotypic behavioural responses (suckling, vomiting, defecation and crying). It represents the primary mechanism that regulates positive and negative feeling states, providing the child with the first value-based map of the world, founded on positive and negative experiences related to food intake, satiation and associated gut responses. For example, the response to sweet taste receptor stimulation of the tongue in the newborn is a stereotypically organized reflex response also seen in decerebrate rats<sup>98</sup> and in anencephalic infants<sup>99</sup>. The concept of a body-based value map is closely related to the 'somatic marker' hypothesis or to the concept of interoceptive memories (BOX 2). For example, the close relationship between sucrose ingestion and the resulting activation of sweet taste receptors on the tongue and central release of opioids (important mediators of homeostasis and well being), has been shown both in rodents<sup>100</sup> and in children<sup>101</sup>. This basic gut-based neurochemical encoding of feeling states is often maintained in the adult but complemented by learned meta-representation of such sensations (FIG. 3). For example, the close relationship of the gastric-based ghrelin signalling system, which plays a prominent part in the generation of the homeostatic emotion of hunger, and the associated craving for food, with the dopamine-based reward system is maintained throughout development, as peripherally released ghrelin can stimulate central dopamine release through vagal and endocrine mechanisms<sup>102</sup>. These initially gut-based responses to appetitive and aversive stimuli may also form the basis for interoceptive brain-based memories of basic emotional states (FIG. 3) that are relevant for complex social emotions, including predictions about the future and intuitive decision making. The fact that predictions and prediction errors are associated with activations of sub-regions of the aINS<sup>103,104</sup>, and that patients with lesion of the OFC, a brain region closely connected to the aINS, are greatly compromised in their ability to make adaptive decisions about the future<sup>105</sup>, provides indirect evidence for such a hypothesis. It has been speculated that a system of specialized cells (von Economo (VEN) cells) that are located in the aINS and share certain receptors and peptides that are rare in other brain regions but abundant in the ENS (including bombesin-like peptides and 5-HT receptor 2B (5HT2B)), may be related in evolutionary terms to gut-brain signals<sup>106</sup>. The proposed close relationship between brain and gut systems is supported by evolutionary, developmental and neurobiological evidence. For example, there is a close relationship between pleasant gut-related emotions (associated with food intake) and social interactions in societies and cultures of human and non-human primates, as well as in lay language. Conversely, there is a close relationship between negative gut-related feeling states (including nausea and abdominal discomfort) and social withdrawal.

The close relationship between gut signalling systems, basic feeling states and meta-representations of these feeling states in basic emotions and behaviours is not limited to nutrient-related signals. Gut-based signalling systems that are related to an insult of gut homeostasis (by chemicals, toxins and inflammation) can easily fit into this general model. This includes the 5HT-5HT<sub>3</sub> receptor-vagal signalling system related to toxin-induced nausea<sup>107</sup>, the cytokine-vagal signalling system<sup>108</sup> that is engaged in the generation of fatigue and sickness behaviour, and the CRF signalling system that is related to the threat of mucosal homeostasis by gut inflammation<sup>109</sup>. The concept of a primordial stress response system that is related to immune activation, which is co-opted into a generalized stress response system for both interoceptive and exteroceptive stressors has previously been suggested<sup>110</sup>. Given the magnitude and complexity of gut-based signalling systems, compared to those of other organs, brain-gut interactions have to be considered an important, but largely ignored, component of the neuroscience of emotions.

## Decisions based on gut feelings

As discussed above, the popular statement that somebody has made a decision based on their gut feelings may have an actual neurobiological basis related to brain–gut interactions, and to interoceptive memories related to such interactions. Specifically, evidence from neuroimaging studies in humans has implicated the fronto-insular cortical regions, in particular subregions of the aINS, in intuitive decision making. Intuitive decision making can be defined as the rapid assessment of the probability of a favourable or unfavourable outcome of a planned behaviour in a situation of uncertain outcomes, which is dependent on previous experiences rather than on serial processes of inductive and or deductive reasoning<sup>103,111</sup>. It has been suggested that large spindle shaped neurons that are primarily located in the fronto-insular region of the brain, and present in humans and other mammals with complex social interactions, may be involved in such unique cognitive processes<sup>106</sup>. It remains to be determined if such intuitive decision making is based on an interoceptive map of gut responses, acquired during infancy and refined during development.

## Perturbation of homeostasis

Considerable experimental evidence demonstrates the plasticity of the brain–gut axis during acute and chronic perturbation of homeostasis, including changes in gut to brain signalling that are associated with increased consumption of fat-rich diets<sup>59</sup> and neuroplastic changes in visceral afferent pathways associated with gut inflammation<sup>74,112</sup>.

## Alterations in signalling systems

Extensive evidence supports the involvement of different peripheral signalling systems in perturbations of gut to brain signalling. For example, when intestinal enterochromaffin cells are activated by potentially noxious stimuli, such as food-related or bacterial toxins (or certain chemotherapeutic drugs), increased 5-HT release activates 5-HT<sub>3</sub> receptors on myenteric IPANs as well as on vagal afferents, resulting not only in exaggerated intestinal motor and secretory reflexes but also in the activation of brain regions receiving vagal input. Activation of this enterochromaffin cell–vagal afferent pathway by harmful substances, including certain pathogens, is associated with the subjective sensations of discomfort and nausea, and may be accompanied by vomiting. Together, the peripheral and central responses initiate a complex adaptive response aimed at expelling the offending agent.

Another well-characterized example of a brain–gut signalling system that is engaged during perturbation of homeostasis is the CRF signalling system. Central release of CRF during experimental stress, or central injection of this peptide in unstressed animals produces a distinct, stereotypic pattern of stress-induced gastrointestinal motor and secretory responses that are associated with anxiety-like behaviours<sup>23</sup>. As the stress-induced gut response can be prevented by either peripheral or central CRF1 receptor antagonism, both peripheral and central CRF–CRF receptor 1 (CRFR1) signalling systems seem to be involved. The slowing or reversal of gastric motility and the acceleration of distal bowel motility and secretion result in the reduced exposure of the gastrointestinal tract to luminal factors, reducing the metabolic demand generated by gut-related motor, secretory and absorptive function, and shifting blood flow to the cardiovascular and skeletomotor system for the fight and flight response.

Inflammatory changes in the viscera, including the liver and the gut, can result in cytokine-mediated vagal activation of the hypothalamus and related limbic brain regions, resulting in hyperalgesia and social withdrawal referred to as ‘sickness behaviour’<sup>113</sup>. Additional non-vagal mechanisms by which peripheral cytokines can reach the CNS have been reported<sup>114,115</sup>. Mucosal release of cytokines and other inflammatory mediators including bradykinin, histamine and proteases can also result in sensitization and activation of spinal

afferent terminals, resulting in visceral hyperalgesia. Multiple effects of such inflammation or toxin-induced changes in afferent terminals have been reported, including release of neuropeptides such as substance P and calcitonin gene-related peptide from afferent nerve terminals<sup>116</sup>, development of mechanosensitivity of previously unresponsive afferents and development of peripheral and central sensitization, resulting in greatly increased signalling of visceral afferent information to the CNS<sup>56</sup>.

### Brain–gut interactions in chronic disease

Even though dysregulations of the brain–gut axis have been implicated in several chronic disorders, convincing evidence to support such a pathophysiological role is limited to several chronic gastrointestinal disorders (including so-called functional gastrointestinal disorders (FGIDs) and inflammatory bowel diseases (IBDs)) and to eating disorders, particularly obesity<sup>117</sup> and anorexia nervosa<sup>89</sup>.

FGIDs comprise a heterogeneous group of disorders characterized by chronically recurring pain and discomfort that is conveyed to different regions of the gastrointestinal tract. They are often associated with alterations in gastrointestinal function, and characterized by the absence of any agreed upon biological, physiological or anatomical abnormalities that would explain the symptoms<sup>118</sup>. In the majority of patients, increased levels of anxiety are present, and co-morbid diagnoses of anxiety and depression are seen in up to 60% of patients. Based on symptom criteria, but in the absence of any syndrome-specific pathophysiological findings, up to 40 different functional gastrointestinal syndromes have been postulated, including functional chest pain of presumed oesophageal origin, functional dyspepsia and irritable bowel syndrome (IBS)<sup>118,119</sup>. A characteristic feature of all of these disorders is the frequent overlap of symptoms belonging to one syndrome with one or several others in the same patient, concurrently or at different life periods. One of the most common and best-studied syndromes is IBS<sup>120,121</sup>. Despite intense clinical and preclinical research efforts during the past 20 years, no single biological abnormality has been identified that would explain the symptoms of chronically recurring pain and discomfort in all of the patients meeting current symptom-based criteria. Based on animal models and on studies performed in human patient populations, peripheral mechanisms that have been implicated include long lasting sensitization of primary afferent pathways innervating the intestine, altered microflora–epithelial–immune interactions (including alterations in mast cell regulation, increased epithelial permeability and altered gut microflora)<sup>122</sup>. Perceptual hypersensitivity to experimental and physiological gut signals is common among patients with IBS<sup>123</sup>, but it remains unclear if this reflects an increased interoceptive signal from the gut, a central amplification mechanism related to altered cognitive and affective modulation of the perception of normal or even reduced gut signals. Based on a growing number of brain imaging studies in patients with IBS, alterations in brain responses to controlled gut distension as well as to expectation of such gut discomfort have been reported<sup>9</sup>. In a recent meta-analysis of 18 studies comparing brain responses to rectal distension in patients with IBS and healthy control subjects, patients with IBS showed greater activation of brain regions that are involved in the processing of visceral afferent information, in emotional arousal and in cortical modulation. Recent evidence suggests the presence of regional cortical thickness changes in corresponding brain regions<sup>41,124</sup>. It remains to be determined whether the observed functional and structural brain changes are secondary to chronically increased gut to brain signalling and/or whether they are primary brain-based abnormalities that play a part in pain amplification and altered autonomic regulation of gut function. An important role of central dysregulation in IBS pathophysiology is suggested by the common co-morbidity with other functional gastrointestinal and non-gastrointestinal syndromes (including, but not limited to, interstitial cystitis or painful bladder syndrome, fibromyalgia and temporomandibular disorder), by the common affective co-morbidity, the prominent

role of stress in symptom onset or exacerbation, the responsiveness of these syndromes to centrally targeted therapies (including tricyclic antidepressants and cognitive behavioural therapies) and the fact that the first onset of abdominal pain symptoms in children is often preceded by increased anxiety<sup>125</sup>.

### **Inflammatory gastrointestinal disorders**

Despite a wealth of preclinical data on the acute and persistent effects of gut inflammation on intrinsic and extrinsic gut innervation (reviewed in REFS 56,74,112,126), the role of chronic intestinal inflammation on symptoms and brain responses in human patients with inflammatory bowel diseases (IBDs), such as ulcerative colitis and Crohn's disease, or gut infections are poorly understood. Surprisingly, abdominal pain or persistent visceral hyperalgesia to experimental stimuli is not a prominent feature in most patients with an uncomplicated IBD, and only a small percentage of patients with IBD or with infectious gastroenteritis, experience persistent pain once the inflammation has resolved<sup>127</sup>. The effective engagement of peripheral<sup>128</sup> as well as central<sup>129</sup> pain inhibition systems have been proposed as plausible mechanisms for this paucity of pain symptoms in the majority of patients with IBD. Similarly to IBS, it remains to be determined whether commonly reported co-morbidity with anxiety and depression<sup>130,131</sup> is a consequence of the primary disease, plays a part in the chronicity of symptoms or is a vulnerability factor. Evidence from animal models supports a modulating role of the brain on gut inflammation<sup>132</sup>.

### **Eating disorders**

Disturbances in ingestive behaviour are common in industrialized societies and both obesity and anorexia nervosa are major health problems in terms of morbidity, mortality and healthcare costs<sup>133,134</sup>. Although obese subjects eat beyond their caloric needs, with food intake driven primarily by the craving for the hedonic value of food, despite the known negative health consequences<sup>117</sup>, individuals with anorexia nervosa have an ego-syntonic resistance to eating and an obsession with weight loss, despite a preoccupation with food and eating rituals<sup>89</sup>.

Despite the extensive progress in the understanding of peripheral and central contributions of altered brain– gut interactions in obesity (reviewed in REFS 117,135), attempts at pharmacological treatments have been disappointing and various gastric surgical procedures are currently the most effective treatment<sup>136</sup>. Even though the current obesity epidemic is likely to be multifactorial, a shift in the balance between hedonic and homeostatic regulation of food intake, a compromised effectiveness of cortical control mechanisms and changes in peripheral encoding mechanisms in the gut seem to play an important part. For example, diet-induced down-regulation of satiety-inducing gut to brain signaling mechanisms have been reported, including leptin and insulin resistance and alterations in the normal cholecystokinin-dependent molecular switching mechanisms in vagal afferents during food intake from an orexigenic to an anorexigenic phenotype<sup>137</sup>. At the same time, considerable evidence for alterations in the regulation of the hedonic aspects of food intake have been reported, consistent with the concept of food addiction<sup>135</sup>. Brain imaging findings in humans who are obese are consistent with an enhanced sensitivity of the reward circuitry to food cues (for example, viewing images of high caloric food) that predict reward, but a decreased sensitivity to the rewarding effects of actual food ingestion (interoceptive stimuli) in dopaminergic pathways. As several of the gut peptides are able to stimulate the central dopamine system (including ghrelin, leptin and insulin), the observed changes in the central reward circuits may in part be due to diet-related neuroplastic alterations in peripheral encoding mechanisms and the reduced responsiveness of interoceptive signalling mechanisms from the gut.

The alterations in the central regulation of food intake in anorexia nervosa have recently been reviewed<sup>89</sup>. Despite the behavioural abnormalities, and more recently the brain abnormalities, that have been reported in patients, little is known about possible alterations in gut to brain signalling mechanisms and in particular about possible alterations in vagally mediated neu-ropeptide signalling to the brain. However, the fact that patients with anorexia nervosa showed reduced responses in homeostatic afferent brain regions (the aINS and ACC) in response to oral sucrose administration and that no correlation between activity in these regions and self-ratings of the pleasantness of the administered sucrose was observed, suggest that the influence of interoceptive signals from the gastrointestinal tract on ingestive behaviour is reduced.

### Proposed disease model for altered brain–gut interactions

Analogous to cognitive models of anxiety<sup>138–140</sup> and based on the central role of ineffective coping strategies (such as catastrophizing or hypervigilance) and anxiety in determining symptom severity in several functional gastrointestinal disorders, a central role of altered appraisal and prediction error has been proposed<sup>125</sup> (FIG. 4). This model posits a mismatch between the actual interoceptive image of the digestive system represented in the INS and the predicted state, based on a negatively valenced interoceptive memory of such experiences. The mismatch results in hypervigilance towards gut-related sensations and emotional arousal, engagement of ANS responses and possibly enhanced sensory perception. In the case of obesity and food addiction, the proposed model postulates a mismatch between the expected reward and the actual delivery of reward. This mismatch will promote compulsive overeating as an attempt to achieve the expected level of reward<sup>135</sup>. The mechanisms that underlie the failure to correct the mismatch by updating the predicted state are unclear but have been discussed in the context of learning theory and reinforcement learning<sup>141,142</sup>.

### Conclusions and future directions

Tremendous progress has been made in our understanding of the bidirectional crosstalk between the brain and the digestive system. This includes the remarkable success in mapping the functional neuroanatomy of the ENS, in our understanding of how the brain modulates these ENS circuits and gut functions, and in unravelling the complexity of gut to brain signalling through multiple parallel but interacting communication channels. Alterations in brain–gut interactions that are associated with gut inflammation, chronic abdominal pain syndromes, eating disorders and psychosocial stressors have been characterized, even though the development of effective therapies for any of these conditions has been disappointing. By contrast, many aspects of gut to brain signalling that are addressed in this article, in particular the role of this signalling in emotional and cognitive function, remain speculative at this point but may guide future research. Several important questions remain to be addressed. What role does gut to brain signalling have in brain development during early life and what part does it play in adults? For example, does gut to brain signalling early in life influence the development of adult ingestive behaviour, visceral pain sensitivity, mood and affect, interoceptive memory and intuitive decision making? What roles do taste receptors and the microflora play in this interoceptive signalling? Does mucosally driven vagal input to the brain play a part in brain development? Is there a role for alterations in microbial signalling to the gut and in gut signalling to the brain in the pathophysiology of anxiety and depression, as suggested by the common overlap of several brain–gut disorders with disorders of mood and affect? And what part do interoceptive memories and prediction errors play in human disease states? One of the great advantages in addressing these questions is the fact that most of these hypotheses can be studied in human patients, from whom stool samples for metagenomic sequencing of the microflora and gut tissue for analysis of signalling mechanisms can be obtained relatively

easily by endoscopic biopsies, and brain structure, function and signalling can be studied with non-invasive neuroimaging techniques.

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## Abbreviations

### **Enteric nervous system**

Ganglionated plexus of neurons that is located between the layers of the gut. These neurons, which are equal in number to those in the spinal cord, are able to regulate basic gut functions, such as the peristaltic reflex.

### **Emotional motor system**

System of parallel outflows from cortico–limbic–pontine networks that is engaged during distinct homeostatic states. A medial component provides tonic modulation of spinal reflexes

	and a lateral component plays a part in executing distinct regional motor patterns of the viscera through autonomic pathways
<b>Enteroendocrine cell</b>	Specialized epithelial cell that releases secretory granules, containing one or several gut peptides, on the basolateral side (and possibly luminal side) in response to luminal chemical, mechanical and possibly neural stimuli.
<b>Enterochromaffin cell</b>	Specialized epithelial cell that releases secretory granules, containing primarily serotonin, on the basolateral side (and possibly luminal side) in response to luminal chemical, mechanical and possibly neural stimuli.
<b>Intrinsic reflex</b>	Also known as an intramural reflex. A reflex of the enteric nervous system in which the afferents, interneuron and efferent neurons that are involved are all contained within the gut wall.
<b>Intrinsic, primary afferent</b>	Afferent neuron with its cell body contained within the enteric nervous system, that encodes mechanical and paracrine signals.
<b>Commensal bacteria</b>	Refers to the 100 trillion bacteria that live in symbiosis with the gut and make up the intestinal microflora.
<b>Myenteric</b>	Subplexus of the enteric nervous system, which is localized between the circular and longitudinal muscle layer.
<b>Ego-syntonic</b>	Psychological term referring to behaviours, values and feelings that are in harmony with, or acceptable to, the needs and goal of a person or that are consistent with a person's ideal self image. This trait is typically seen in patients with anorexia nervosa.

**Box 1****Evolutionary and developmental aspects of the enteric nervous system**

From an evolutionary standpoint, it is clear that the enteric nervous system (ENS) is not uniquely human or even mammalian: homologues of an ENS are found throughout the animal kingdom, including in insects, snails and marine polyps<sup>143–145</sup>. It has been suggested that the ganglia that form the primitive brains of helminths, and eventually the brains of higher mammals, were derived from the more primitive but homologous enteric nerve circuits<sup>146</sup>. Thus, neural circuitries and transmitter systems that have evolved to assure optimal responses (approach and withdrawal responses) to the challenges presented by our internal — for example, luminal — environment may have been incorporated into the CNS during evolution. Developmentally, the ENS arises from precursor cells that migrate from the neural crest along the vagus to settle and differentiate in the gut<sup>147</sup>. Based on its close bidirectional connections with limbic and autonomic regions of the brain<sup>148</sup>, the ENS can be viewed as a peripheral extension of the limbic system into the gut, where it is exposed closely to our complex internal environment, including powerful mechanical, chemical and microbial influences. Alternatively, parts of the CNS (in particular, pontine, autonomic and limbic circuits) can be viewed as an encephalized portion of the ENS.

**Box 2****Influential theories of brain–viscera interactions**

The first comprehensive scientific theory of brain–viscera interactions was formulated in the 1880s by William James and Carl Lange and was based on the central concept that stimuli that induce emotions such as fear, anger or love initially induce changes in visceral function through autonomic nervous system output, and that the afferent feedback of these peripheral changes to the brain is essential in the generation of specific emotional feelings<sup>8</sup>. According to this theory, we feel anxious because we perceive our heart beating faster, because we become aware of our respiration becoming more frequent and shallower, or because we feel ‘butterflies’ in our stomach. In the late 1920s, Walter Cannon challenged the James–Lange theory, postulating that emotional feelings are generated directly by subcortical brain regions rather than from the feedback of situational bodily changes. He proposed that such bodily changes that are associated with emotional states are simply by-products of these brain changes, and that the visceral responses are too slow to play any part in the subjective experience of emotional feelings<sup>7</sup>.

Modern theories of emotion and consciousness that were proposed in the form of the ‘somatic marker’ hypothesis by Antonio Damasio<sup>81</sup>, and the ‘homeostatic emotion’ hypothesis by A. D. Craig<sup>78</sup> have re-emphasized the importance of interoceptive feedback in emotional states and cognitive processes, and have largely overcome the long lasting controversy about the directionality of brain–viscera interactions in the generation of emotions.

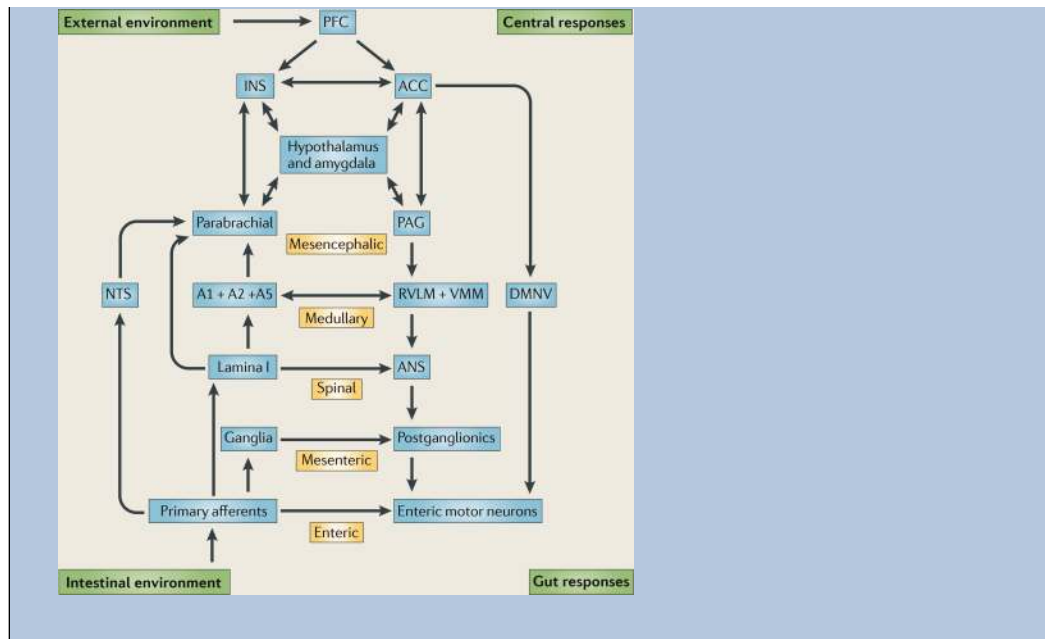
For example, Damasio eloquently proposed that somatic markers (for example, memories of body states associated with previous feeling states) arise from positive or negative emotional feeling states being associated with visceral and other bodily responses (body loops) to certain contextual situations. According to this theory, these body loops, or their meta-representations in the orbitofrontal cortex (OFC), may play a part not only in how somebody feels at a given moment but may also influence future planning and intuitive decision making. For example, according to Damasio, somatic markers may covertly result in “undeliberated inhibition of a response learned previously ... [or] the introduction of a bias in the selection of an aversive or appetitive mode of behavior”<sup>81,149</sup>.

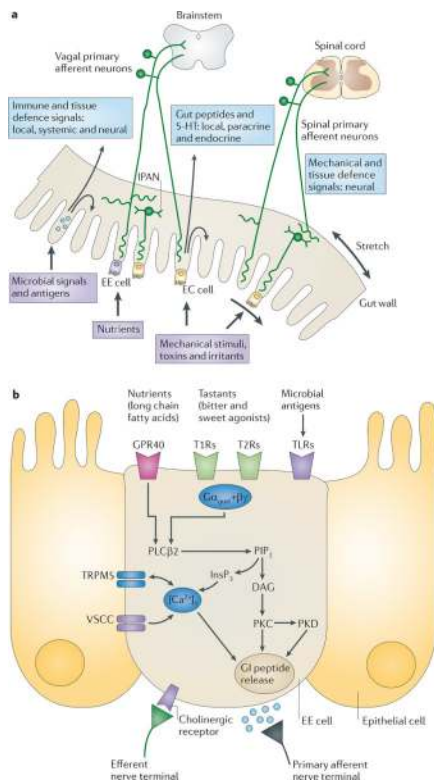
Craig's homeostatic emotion hypothesis postulates that, in humans and anthropoid primates, an image of the homeostatic state of the body is created in the primary interoceptive cortex (dorsal posterior insula) through lamina I spinothalamic and vagal afferent tract projection to a specific thalamocortical relay nucleus in the posterolateral thalamus<sup>77</sup>. This interoceptive cortex contains modality-selective representations of all afferent activity from lamina I (that is, sympathetic afferent input) and from the nucleus tractus solitarius (parasympathetic input). According to Craig, re-representation of the image from the posterior insula to the mid and anterior insula, and modulation by inputs from affective, cognitive and reward-related brain circuits creates a feeling about the homeostatic state of the body in the anterior insula<sup>78,80</sup>. Together with the parallel processed signal in the anterior cingulate cortex (ACC), a homeostatic emotion arises, made up of a feeling dimension represented in the anterior insula and a motivation dimension represented in the ACC<sup>78</sup>.

**Box 3****The brain–gut axis is a system of hierarchical homeostatic reflexes**

The brain–gut axis can be conceptualized as a hierarchy of reflexes, from the reflex circuits contained within the enteric nervous system to the highest reflex loop involving the insular cortex and anterior cingulate cortex (ACC) (see the figure). Whereas many local physiological responses of the intestine to stretch or chemical stimulation (peristaltic, secreto-motor reflex) only involve enteric reflexes (for example, the entire reflex loop is contained within the bowel wall), most gastric reflexes and reflexes involving spatially distinct regions of the gastrointestinal tract (for example, duodeno–gastric feedback pathways in the regulation of gastric emptying) involve mesenteric reflexes (the reflex loop involves mesenteric ganglia) or vago-vagal reflexes (the reflex loop involves the dorsal vagal nucleus). Gut responses to nociceptive stimuli typically involve spinal and supraspinal reflexes, engaging strong emotional and autonomic responses. The engagement of circuits outside of the gut wall integrates interoceptive and exteroceptive information to optimize homeostatic regulation of intestinal function. Pontomedullary nuclei (including the midline raphe nuclei and the locus coeruleus complex) exert a tonic inhibitory influence on the gain of these reflexes, whereas top-down influences that originate in prefrontal regions can regulate sympathetic and vagal outputs as well as modulate the dorsal horn through descending influences. Top-down influences are engaged in responses to environmental stimuli (threat and safety) and in the recall of memories, including interoceptive memories. The conscious awareness of interoceptive images that is generated either by signals from the gut or by the recall of interoceptive memories of such signals is associated with the perception of emotional feelings, including pain, disgust or well being. The nucleus tractus solitarius (NTS) is the vagal relay nucleus in the medulla that receives input from function-specific vagal afferents and signals through interneurons to the vagal premotor neurons in the dorsal motor nucleus of vagus (DMNV). The NTS and DMNV make up the dorsal vagal complex (DVC). A1, A2 and A5 are medullary catecholaminergic nuclei to autonomic effector regions in the rostro–ventro–lateral medulla (RVLM) and ventromedial medulla (VMM). The periaqueductal grey (PAG) receives inputs from the parabrachial nucleus and also from forebrain regions, including the hypothalamus, amygdala and anterior cingulate cortex, and the prefrontal cortex (PFC). It is organized into function-specific columns. The PFC provides the highest cortical modulatory input to the insula (INS) and ACC. ANS, autonomic nervous system.





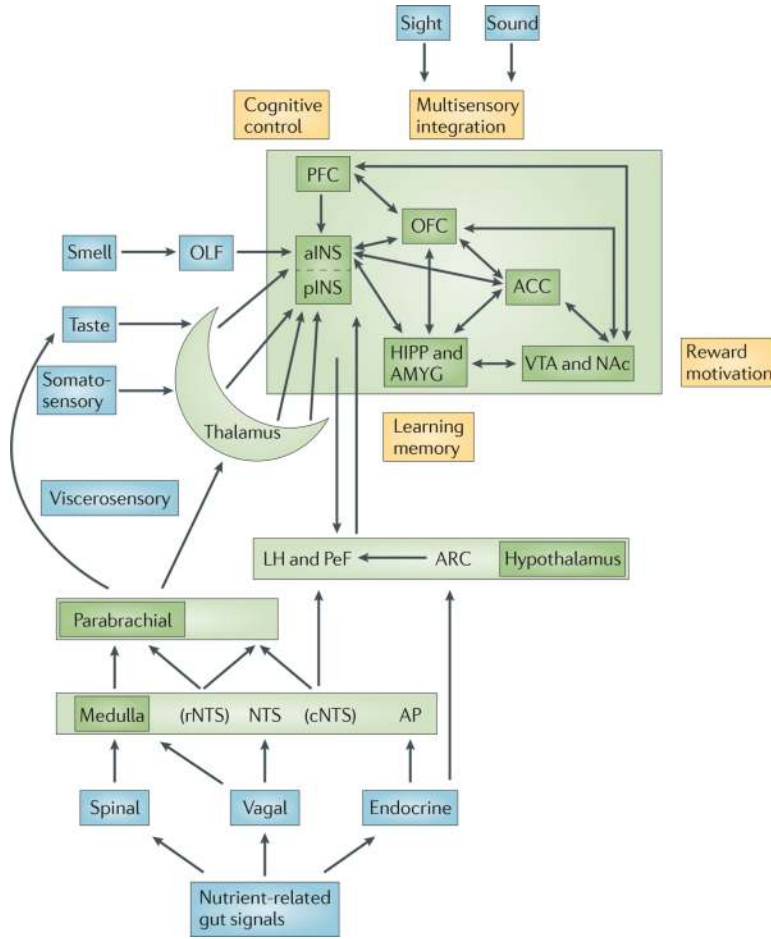


**Figure 1. Gut to brain communication**

**a** | Endocrine, immune and neuronal afferent signalling from the gut to the CNS.

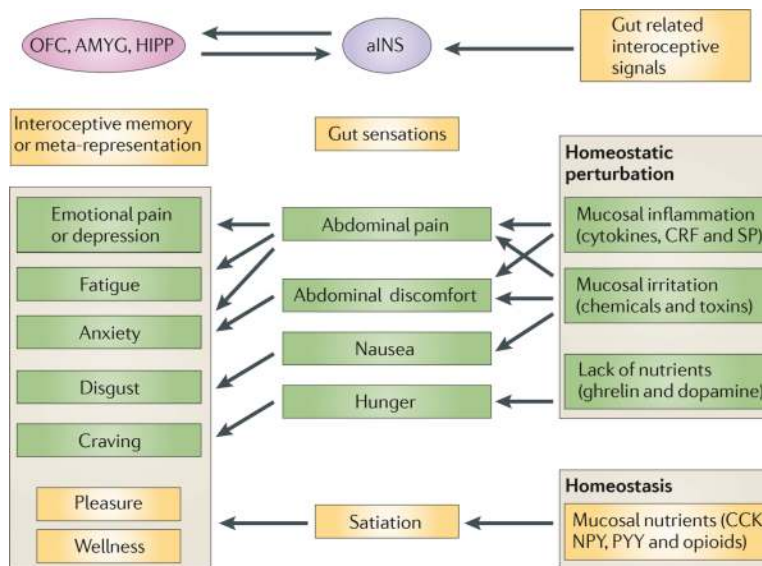
Information about luminal factors and conditions of the gut are signalled through extrinsic vagal and spinal afferents to the brain stem and spinal cord, respectively. Mechanical stimuli (stretch, pressure, distortion and shearing forces) can activate spinal, vagal and intrinsic primary afferents (IPANs) directly, without intermediary cells such as the enteroendocrine (EE) cells. Although no synaptic connections have been found between IPANs and extrinsic afferents, the latter form networks around myenteric ganglia (intraganglionic laminar endings<sup>27</sup>), many of which receive synaptic input from IPANs. Signalling molecules (including proteases, histamine, serotonin and cytokines) that are produced by immune cells in Peyer's patches and within the gut epithelium can activate their respective receptors on vagal and spinal afferents. Similarly, neuropeptides and hormones (gut peptides) that are released from EE cells in response to other luminal factors, such as nutrients, toxins or antigens, can act both in an endocrine fashion, reaching targets in the brain (area postrema, dorsal vagal complex and hypothalamus), and through receptor activation on spinal and vagal afferents, in a paracrine fashion. Enterochromaffin (EC) cells signal to both IPANs and vagal afferents. **b** | Encoding of multiple luminal signals by EE cells. Different classes of EE cells are interspersed between gut epithelial cells throughout the gastrointestinal tract. Upon luminal stimulation (or upon activation by postganglionic sympathetic or vagal nerves), these cells can release up to 20 different gut peptides from their basolateral (and possibly luminal) surface. Released peptides can activate closely adjacent vagal afferent nerve terminals in a paracrine fashion, or when released into the circulation they can exert an endocrine effect, signalling to various sites in the brain and other parts of the gastrointestinal tract. Different types of receptors have been identified on the luminal side of EE cells, including G protein-coupled taste receptors (GPCRs) for sweet and bitter tastants, GPCRs that are responsive to fatty acids and toll-like receptors (TLRs). The intestinal taste receptors that are shown are coupled to a specific G protein subunit, gustducin (G<sub>gust</sub>),

and receptor-induced increases in intracellular calcium result in peptide release from the basolateral membrane.  $[Ca^{2+}]_i$ , intracellular calcium concentration; DAG, diacylglycerol; GI peptide, gastrointestinal peptide; GPR40, G protein-coupled receptor 40;  $InsP_3$ , Inositol-1,4,5-trisphosphate;  $PIP_2$ , aquaporin PIP2 member; PKC, protein kinase C; PLC  $\beta$ , phospholipase C  $\beta$ ; T1R, taste receptor type 1 member; TRPM5, transient receptor potential cation channel subfamily M member 5 (specifically linked to taste receptor signalling); VSCC, voltage-sensitive  $Ca^{2+}$  channel.

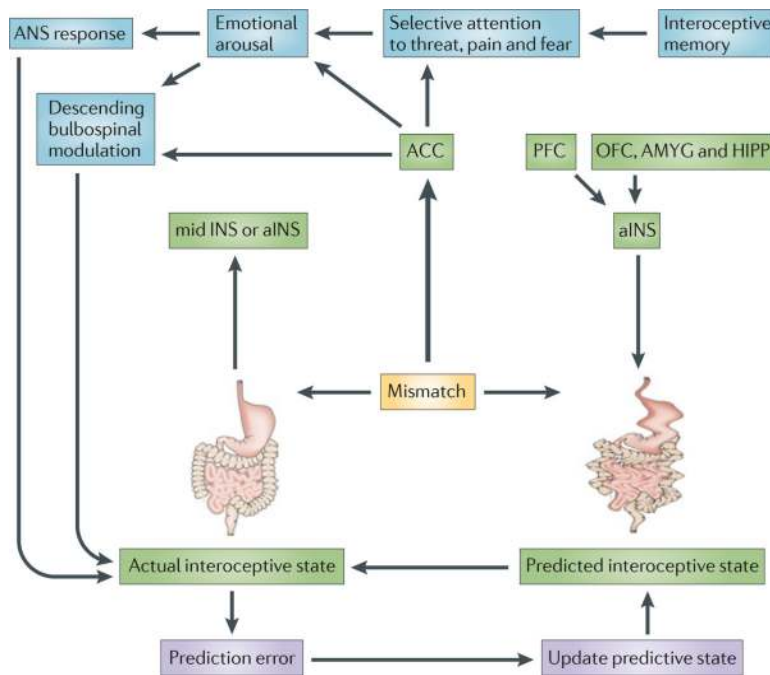


**Figure 2. Gut–brain signalling related to food intake**

Nutrient-related signals reach the CNS through spinal, vagal and endocrine signalling pathways. Endocrine signalling of gut peptides that are released into the systemic circulation reach the dorsal vagal complex through the area postrema where they modulate the transmission of afferent vagal signals to the dorsal motor nucleus. These gut peptides also reach specialized neurons within the hypothalamus. Paracrine signals activate function-specific vagal afferent fibres that ultimately signal to subregions of the anterior insula (aINS). The sensory aspect of taste is primarily encoded in the aINS, but the multimodal integration of satiety signals with the sensory properties of food (including its flavour, palatability and reward value) as well as the context of food intake (including food related visual and auditory signals) occurs in the orbitofrontal cortex (OFC). Further integration with inputs from the reward system and with interoceptive memories of previous food ingestion generates a multidimensional food-related experience that ultimately determines ingestive behaviour. Prefrontal regions exert cognitive control over ingestive behaviours. Learning about food-related experiences and the formation of interoceptive memories is an important aspect of the cortical circuitry that is involved in this process. ACC, anterior cingulate cortex; AP, area postrema; ARC, arcuate nucleus; cNTS, caudal NTS; HIPP, hippocampus; LH, lateral hypothalamus; NAc, nucleus accumbens; NTS, nucleus tractus solitarius; PeF, perifornical hypothalamus; OLF, olfaction; PFC, prefrontal cortex; rNTS, rostral NTS; VTA, ventral tegmental area.



**Figure 3. Gut signalling systems, gut sensations and meta-representations of such sensations**  
 Gastrointestinal interoceptive signals to the brain play a key part in the maintenance of homeostasis under resting conditions. Whereas signalling under homeostatic conditions is generally associated with pleasant sensations (satiety), signalling under non-homeostatic conditions is typically associated with aversive sensations (pain and nausea). Perception of these signals as gut sensations is associated with activation of the anterior insula (aINS) and the orbitofrontal cortex (OFC), where gustatory, olfactory and viscerosensory signals are integrated and where they are modulated by affective and cognitive input. The generation of such gut sensations can lead to the formation of interoceptive memories of the multidimensional experience associated with the sensation (such as the feeling of nausea associated with a disgusting experience or a particular taste with a pleasant social experience). The interoceptive memory is encoded within a distributed network involving the aINS, OFC, the amygdala (AMYG) and hippocampus (HIPP). Often, environmental stimuli that are unrelated to the gut, such as an image, sound or smell, may trigger the recall of these feeling states as meta-representations of the original gut-triggered sensation. The experience of such memories can result in the engagement of similar brain regions (aINS, anterior cingulate cortex (ACC), OFC and amygdala) in the absence of gut signalling. This recall may be the basis for feelings such as disgust, craving or emotional pain, in the absence of any interoceptive input from the gut, and may occur when watching another person in pain, when hearing about a disgusting experience or when viewing images of rich foods. Recall of such memories without conscious awareness may bias behavioural responses<sup>149</sup>, and such behavioural biases may be a reflection of an internal value map based on ‘gut feelings’. 5-HT, 5-hydroxytryptamine (serotonin); CCK, cholecystokinin; CRF, corticotropin releasing factor; NPY, neuropeptide Y; PYY, peptide YY; SP, substance P.



**Figure 4. Interoceptive memory and prediction error in chronic disease**

According to a theory proposed by Paulus and Stein<sup>138</sup>, the mismatch of actual interoceptive input reaching the anterior insula (aINS) through body to brain signalling, with a falsely predicted interoceptive state (from interoceptive memory and/or influences on the aINS from prefrontal and limbic influences) results in the engagement of anterior cingulate cortex (ACC). Mismatch-related ACC activation is associated with emotional arousal, increased sympathetic and sacral parasympathetic activity and engagement of descending bulbospinal sensory facilitation systems, and may be associated with a conscious feeling of anxiety and worry. When adapted to functional gastrointestinal disorders<sup>125</sup>, the autonomic response to the mismatch is likely to change the state of the gut through modulation of multiple target cells (for example, activation of motor and secretory activity in the distal colon that is associated with stress), which in turn is likely to produce altered interoceptive feedback to the INS. The failure to correct the prediction error by updating the predictive state (as would be expected in a healthy individual) results in chronicity of symptoms through chronic functional and structural dysregulation of the brain–gut axis. Mismatches between actual interoceptive input from nutrient related signals from the gut and interoceptive memories of hedonic experiences of food intake have been implicated in obesity<sup>135</sup>. ACC, anterior cingulate cortex; AMYG amygdala; ANS, autonomic nervous system; HIPP, hippocampus; OFC orbitofrontal cortex; PFC prefrontal cortex.