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Enterochromaffin cells are gut chemosensors that couple to sensory neural pathways

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Summary

Dietary, microbial, and inflammatory factors modulate the gut-brain axis and influence physiological processes ranging from metabolism to cognition. The gut epithelium is a principle site for detecting such agents, but precisely how it communicates with neural elements is poorly understood. Serotonergic enterochromaffin (EC) cells are proposed to fulfill this role by acting as chemosensors, but understanding how these rare and unique cell types transduce chemosensory information to the nervous system has been hampered by their paucity and inaccessibility to single cell measurements. Here, we circumvent this limitation by exploiting cultured intestinal organoids together with single cell measurements to elucidate intrinsic biophysical, pharmacological, and genetic properties of EC cells. We show that EC cells express specific chemosensory receptors, are electrically excitable, and modulate serotonininsensitive primary afferent nerve fibers via synaptic connections, enabling them to detect and transduce environmental, metabolic, and homeostatic information from the gut directly to the nervous system.

Graphical abstract

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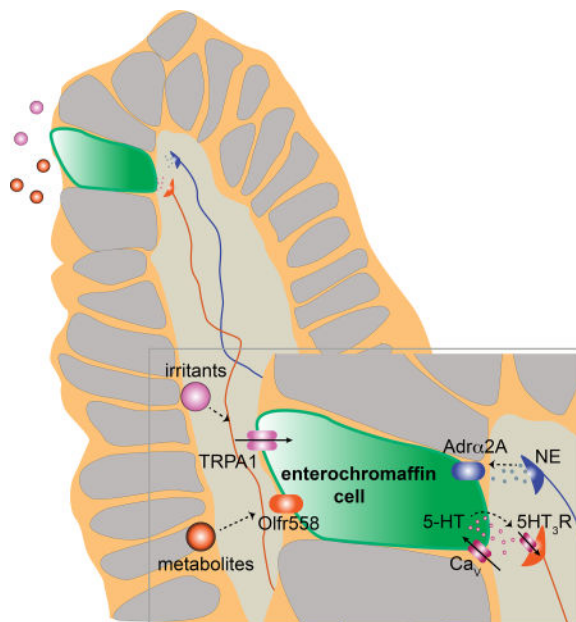
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Use of organoids to characterize rare chemosensory cells in the gut elucidated their biophysical, pharmacological and genetic properties, and showed that they communicate directly with neural sensory pathways.

Introduction

The gut epithelium forms one of the largest exposed surfaces of the human body, representing a unique interface for integrating environmental information with physiologic signals from nervous, immune, and vascular systems (Furness et al., 2013; Ohman et al., 2015). Dietary nutrients and irritants, microbiota products, and inflammatory agents have been proposed to act on the gut epithelium to modulate downstream signaling pathways controlling digestion, immunity, metabolism, and pain (Brierley and Linden, 2014; Furness et al., 2013; Gribble and Reimann, 2016). Hormone-producing epithelial endocrine cells within the gut form anatomical connections with neurons (Bohorquez et al., 2015), consistent with the idea that the epithelium participates in neural monitoring of the gut environment. Despite growing interest in the gut-neural axis, relatively little is known about molecular mechanisms underlying chemosensory transduction by the gut epithelium, or how this information is transmitted to the nervous system.

The enterochromaffin (EC) cell is an intestinal endocrine cell subtype that represents one of the major proposed epithelial chemosensors. Though relatively rare (<1% of total intestinal epithelia), these unique cells produce >90% of the body's serotonin and have been suggested to affect a variety of physiological and pathophysiological states, such as gastrointestinal (GI) motility and secretion, nausea, and visceral hypersensitivity (Gershon, 2013; Mawe and Hoffman, 2013). Previous studies suggest that EC cells express sensory receptors whose activation promotes release of 5-HT with consequent stimulation of gut contractility. While informative, these initial insights have generally been gleaned from model endocrine tumor

cell lines or *ex vivo* whole tissue preparations (Braun et al., 2007; Doihara et al., 2009; Fukumoto et al., 2003; Hagbom et al., 2011; Kim et al., 2001; Nozawa et al., 2009), leaving fundamental questions unanswered. For example, what stimuli or downstream signaling pathways promote serotonin release from EC cells, and how do EC cells communicate with the nervous system?

To address such questions it is necessary to directly interrogate single EC cells, but this is technically challenging given their relatively small size (<10 μ m) and paucity. We therefore generated intestinal organoids in which EC cells are genetically tagged to facilitate their detailed physiologic, pharmacologic, and molecular characterization in a ‘native’ environment. We show that EC cells are electrically excitable because they express functional voltage-gated sodium (Na⁺) and calcium (Ca²⁺) channels, akin to other primary sensory cells. Moreover, EC cells use specific receptors and signal transduction pathways to detect relevant stimuli: the transient receptor potential A1 (TRPA1) ion channel serves as an irritant receptor, olfactory receptor 558 (Olfr558) serves as a microbial metabolite sensor, and an α 2A adrenergic receptor (Adra2A)-TRPC4 channel signaling cascade detects stress response-related catecholamines. These sensory transduction pathways stimulate P/Q-type voltage-gated Ca²⁺ channels to control serotonin release onto 5HT₃ receptor-expressing primary afferent nerve fibers that extend into intestinal villi and engage in synaptic-like contacts with EC cells. Our findings establish EC cells as specialized, polymodal stimulus detectors that constitute a direct line of communication between the gut epithelium and specific primary afferent nerve fibers. This signaling system represents a key pathway for monitoring GI commensal or infectious microbes, injury, or other changes to the luminal environment that stimulate physiologic responses such as emesis, motility, and visceral pain.

Results

EC cells are electrically excitable

We used mice expressing a genetically encoded fluorescent protein (hrGFP) under control of the *chromogranin A* (*ChgA*) promoter to selectively label EC cells for single-cell characterization (Fig. 1A). Consistent with previous results (Engelstoft et al., 2015), we found that ChgA-GFP co-localized with serotonin, but not with lysozyme or GLP-1, which mark Paneth or peptide hormone-producing enteroendocrine cells, respectively (Fig. 1A and S1A–C). Furthermore, transcriptional profiling of ChgA-GFP⁺ cells revealed enrichment of EC cell markers, but not markers for hormone-producing enteroendocrine cells (Fig. S1D). We used whole-cell patch-clamp recording to analyze the electrophysiological properties of single EC cells from native intestinal tissue or intestinal organoids derived from these animals (Fig. 1B, C). We first observed a large outward current in response to voltage ramps that was blocked by tetraethylammonium (TEA⁺) (Fig. 1B), indicating that the current is carried by K⁺ channels, several subtypes of which were expressed at high levels in EC cells (Fig. S2A). Acute TEA⁺ treatment revealed small voltage-activated inward currents that we further characterized in the presence of intracellular Cs⁺ to block K⁺ channels (Fig. 1B). Under these conditions, voltage-steps elicited rapidly-inactivating tetrodotoxin-sensitive inward currents in EC cells from intestinal organoids or native intestine, suggesting the presence of voltage-gated sodium (Na_v) channels (Fig. 1C, D). Steady-state activation and

inactivation properties of this current were consistent with properties of the $\text{Na}_V1.3$ subtype of voltage-gated Na^+ channels (Catterall et al., 2005a), whose transcripts were highly enriched in EC cells (Fig. 1E and S2B). Furthermore, we found that EC cells are electrically excitable, producing tetrodotoxin-sensitive action potentials (Fig. 1F).

In addition to the transient Na_V current, we observed a small voltage-activated slowly-inactivating current (Fig. S2C). We reasoned that voltage-gated Ca^{2+} channels (Ca_V) might contribute to this current because of its slow inactivation kinetics and the fact that Ca_V channels have been previously implicated in EC cell function (Racke and Schworer, 1993; Raghupathi et al., 2013). Indeed, when intestinal epithelial cells were depolarized with high extracellular K^+ , we observed large increases in intracellular Ca^{2+} predominantly within EC cells (Fig. 1G, H). Moreover, depolarization-evoked Ca^{2+} responses were inhibited by ω -agatoxin IVA, which blocks presynaptic P/Q-type Ca_V channels (Catterall et al., 2005b) (Fig. 1H, I). A subset of EC cells were basally active, exhibiting spontaneous Ca^{2+} bursts that were attenuated by tetrodotoxin or blocked by ω -agatoxin IVA (Fig. S2D). Consistent with this pharmacological profile, P/Q-type $\text{Ca}_V2.1$ channel transcripts were selectively expressed in EC cells (Fig. 1J). Transcripts for the low voltage threshold T-type channel, $\text{Ca}_V3.2$, were also highly enriched in EC cells. Perhaps not surprisingly, a T-type Ca_V channel inhibitor did not significantly affect K^+ -induced responses (Fig. 1I), consistent with the fact that $\text{Ca}_V3.2$ often functions in amplifying signals from low threshold stimuli or regulating membrane potential (Catterall et al., 2005b). In summary, our results show that EC cells express voltage-gated ion channels, rendering them electrically excitable - a hallmark of sensory cell types.

EC cells are polymodal chemosensors

To ask if EC cells serve a specific chemosensory role, we screened 30 potential agonists, focusing on compounds known to be present in the gut, including microbial products, irritants and inflammatory agents, and neurotransmitters. Among these, only allyl isothiocyanate (AITC), isovalerate, and the catecholamines dopamine, epinephrine, or norepinephrine specifically and consistently activated EC cells (Fig. 2A). Additionally, isobutyrate and butyrate elicited small, but consistent responses (Fig. 2A). Each of these compounds evoked Ca^{2+} transients, or in some cases oscillatory responses resembling the Na_V and Ca_V -dependent bursting activity that we observed in subsets of EC cells (Fig. S2D). Remarkably, this activity profile is directly relevant to the etiology of GI inflammation. For instance, AITC, the pungent agent in wasabi and other mustard plants, is representative of a class of reactive chemical irritants that elicit cutaneous and visceral inflammatory pain (Bautista et al., 2006; Brierley et al., 2009). Isovalerate, isobutyrate, and butyrate are volatile fatty acid fermentation products produced by gut microbiota that modulate serotonin biosynthesis (Yano et al., 2015) and are linked to several pathophysiological states (Koh et al., 2016). Lastly, homeostatic regulation of sympathetic signaling increases norepinephrine levels in the gut following infection, injury, and other types of GI stress (Gabanyi et al., 2016). Together, these results support the idea that EC cells serve as sentinels of noxious chemical stimuli or other insults affecting the GI tract.

Our EC cell transcriptional profile showed that sensory receptors are among the most enriched transcripts, with relative expression rivaling the EC cell marker tryptophan hydroxylase 1 (*Tph1*) or the pore-forming subunit of Na_v1.3 (*Scn3a*) (Fig. 2B). Moreover, three of the most enriched sensory receptors are known to be involved in the detection of molecules identified in our screen. For example, transcripts encoding the AITC receptor, TRPA1, were enriched in EC cells (Fig. 2B), consistent with previous histological analysis of intestinal tissue sections (Nozawa et al., 2009). Transcripts encoding olfactory receptors have been previously detected in EC cells (Braun et al., 2007) and our analysis similarly identified Olfr558 as a prevalent sensory receptor in these cells (Fig. 2B). Indeed, orthologues of Olfr558 are widely expressed outside of the olfactory epithelium, including in endocrine cells of the GI tract, where expression is modulated by microbiota (Flegel et al., 2013; Priori et al., 2015). Consistent with our pharmacological screen, isovalerate has been identified as a putative agonist for Olfr558 (Audouze et al., 2014). Finally, the receptor-operated ion channel, TRPC4, is also preferentially expressed in EC cells (Fig. 2B), where it functions as a key component of the catecholamine signal transduction pathway (see below). Transcripts encoding these sensory receptors and transducers were also expressed in EC cells from native intestine (Fig. S3) (Nozawa et al., 2009). Thus, EC cells are polymodal chemosensors that are molecularly tuned to detect physiologically relevant stimuli.

Mechanism of EC cell activation by chemical irritants or microbial metabolites

Expression of TRPA1 channels is sufficient to account for sensitivity of EC cells to AITC and other electrophilic irritants. These responses were blocked by the selective antagonist A967079, further substantiating expression of functional TRPA1 channels by EC cells (Fig. 2C).

To determine whether Olfr558 exhibits a pharmacological profile consistent with metabolite sensitivities observed in EC cells, we expressed mouse Olfr558 in HEK293 cells together with chimeric G $\alpha_{olf/15}$, which couples olfactory receptors to intracellular Ca²⁺ release as a proxy for receptor activity (Zhuang and Matsunami, 2008). Isovalerate evoked large Ca²⁺ responses in these cells, but not in G $\alpha_{olf/15}$ -transfected controls lacking Olfr558 (Fig. 2D, E). Consistent with native EC cell responses, isobutyrate or butyrate evoked smaller responses in fewer cells, while propionate or acetate were inactive (Fig. 2D, E). Human Olfr558 exhibited similar metabolite sensitivities (Fig. S4A). In EC cells, isovalerate-evoked responses were abolished by pretreatment with the G $\alpha_{olf/s}$ regulator cholera toxin or the adenylyl cyclase inhibitor SQ22536, but were not perturbed by inhibitors of other signaling cascades (Fig. 2F). Furthermore, responses were not observed in Ca²⁺-free extracellular solution and were significantly reduced by the P/Q-type Ca_v channel inhibitor ω -agatoxin IVA (Fig. 2F). Together, these results suggest that isovalerate activates a G $\alpha_{olf/s}$ -adenylyl cyclase signaling cascade in EC cells (akin to the canonical transduction pathway in olfactory epithelium), promoting Ca²⁺ influx through downstream Ca_v channels.

Finally, to ask if Olfr558 is required for isovalerate-induced responses in EC cells, we used CRISPR/Cas9 to disrupt the Olfr558 gene in ChgA-GFP intestinal organoids (Fig. S4B, C). Doing so abolished isovalerate-evoked responses but did not affect AITC sensitivity, demonstrating that Olfr558 is required for isovalerate signaling in EC cells (Fig. 2F, G and

Fig. S4D). Collectively, our data suggest that EC cells use Olfr558 to detect specific microbial metabolites present in gut and that the receptor is most selective for isovalerate.

Catecholamine signal transduction in EC cells

Gut catecholamine levels, particularly norepinephrine, fluctuate with infection, inflammation, or altered sympathetic tone (Gabanyi et al., 2016). Consistent with a connection to sympathetic signaling and stress, EC cells were most sensitive to epinephrine and norepinephrine, which evoked large EC cell-specific responses (Fig. S5A). Dopamine was ~100-fold less potent, suggesting that an adrenergic receptor(s) underlies catecholamine sensitivity in these cells. Indeed, yohimbine, an α_2 -adrenoreceptor subtype-selective antagonist, inhibited responses to all catecholamines (Fig. 3A,B). Furthermore, EC cells were activated by clonidine, an α_2 -adrenoreceptor-selective agonist, but not by agonists for other adrenoreceptor subtypes (Fig. S5B, C). Among α_2 -adrenoreceptors, only Adra2A was transcriptionally expressed in EC cells, albeit at relatively low levels and in other intestinal epithelial cells (Fig. S5D). However, Adra2A immunoreactivity was specific to EC cells and localized to the basolateral surface, suggesting that protein expression is enriched in EC cells due to preferential translation or enhanced protein stability (Fig. 3C). Tyrosine hydroxylase (TH), a marker for norepinephrine-producing sympathetic fibers, was localized adjacent to EC cells (Fig. 3D), suggesting that sympathetic output can stimulate EC cells by acting on basolateral Adra2A.

How does adrenergic signaling activate EC cells? Inhibition of $G\alpha_i$ by pertussis toxin blocked epinephrine-evoked Ca^{2+} responses (Fig. S5E, F). Furthermore, responses were attenuated in the absence of extracellular Ca^{2+} , but Ca^{2+} influx was not dependent on Na_V or Ca_V channels since neither tetrodotoxin nor ω -agatoxin IVA abolished responses (Fig. 3E and S5G). Another candidate transducer, namely TRPC4, is a Ca^{2+} -permeable channel that can be activated downstream of $G\alpha_i$ -coupled receptors (Jeon et al., 2012) and is highly enriched in EC cells (Fig. 2B). Indeed, the pan-TRPC blocker, 2-aminoethoxydiphenyl borate (2-APB), or the TRPC4-specific inhibitor, ML204, abolished epinephrine-evoked responses, whereas the TRPA1-selective inhibitor, A967079, had no effect (Fig. 3E and S5H). These results suggest that TRPC4 contributes to the Ca^{2+} -permeable conductance stimulated downstream of adrenoreceptor activation in EC cells.

As further evidence that Adra2A and TRPC4 can form a catecholamine-sensitive signaling cascade, we measured large epinephrine-evoked Ca^{2+} responses in HEK293 cells heterologously expressing both proteins, whereas no responses were seen in cells expressing Adra2A or TRPC4 alone, or Adra2A plus TRPC1, TRPC3, or TRPC6 (Fig. S5I, J). We also observed large, ML204-sensitive epinephrine-evoked currents in HEK293 co-expressing Adra2A and TRPC4, but not in cells individually expressing these proteins (Fig. 3F–H). Consistent with our results from EC cells implicating $G\alpha_i$ -dependent signaling, these currents were blocked by pertussis toxin or by co-expression of a dominant-negative $G\alpha_i$ protein. Furthermore, co-expression of a constitutively active $G\alpha_i$ induced ML204-sensitive TRPC4 currents that occluded epinephrine-evoked responses (Fig. 3F, H). Taken together, our results demonstrate that Adra2A and TRPC4 mediate catecholamine sensitivity in EC cells via a $G\alpha_i$ -dependent signaling cascade.

EC cell stimulation promotes Ca_V -dependent serotonin release

EC cells are serotonergic, but given their low abundance, serotonin release has typically been measured in bulk from intact tissue or model endocrine cell lines (Bertrand et al., 2008; Hagbom et al., 2011; Kim et al., 2001; Nozawa et al., 2009; Yano et al., 2015). To measure release from single EC cells directly, we monitored stimulus-evoked changes in cytoplasmic Ca^{2+} in GFP^+ EC cells within intact organoids, while simultaneously measuring whole-cell currents in adjacent biosensor cells expressing a serotonin-gated ion channel ($5\text{-HT}_3\text{R}$) (Fig. 4A). Robust epinephrine-evoked Ca^{2+} responses in EC cells were quickly followed by large $5\text{HT}_3\text{R}$ currents in adjacent biosensor cells (Fig. 4B). Epinephrine-dependent currents were comparable to maximal currents evoked by a saturating concentration of the $5\text{HT}_3\text{R}$ agonist, mCPBG (Fig. 4B). High extracellular K^+ also evoked large Ca^{2+} responses and $5\text{HT}_3\text{R}$ currents, suggesting that EC cell depolarization is sufficient to induce serotonin release (Fig. 4B). Importantly, when biosensor cells were relocated to GFP -negative epithelial cells, neither epinephrine nor K^+ elicited epithelial cell Ca^{2+} responses or $5\text{HT}_3\text{R}$ currents. As expected, direct activation of $5\text{HT}_3\text{R}$ by mCPBG evoked large currents in biosensor cells, but did not elicit Ca^{2+} responses in EC cells (Fig. 4A, B). Together, these data show that epinephrine promotes serotonin release directly and efficaciously from EC cells to produce local effects.

To determine how receptor-mediated signaling drives serotonin release, we perturbed major components of relevant transduction cascades while simultaneously measuring EC and biosensor cell responses. For example, yohimbine or ML204 blocked epinephrine-induced Ca^{2+} responses and $5\text{HT}_3\text{R}$ currents, confirming that the EC catecholamine signal transduction pathway controls serotonin release (Fig. 4B, C, F). Inhibition of Na_V channels with tetrodotoxin had no significant effect, whereas blockade of Ca^{2+} channels by ω -agatoxin IVA reduced Ca^{2+} responses in EC cells only slightly while completely abrogating $5\text{HT}_3\text{R}$ currents in biosensor cells (Fig. 4C, F and Fig. S6A, B). Similar results were seen with AITC (Fig. 4D, F). Isovalerate also evoked EC cell Ca^{2+} responses that correlated with $5\text{HT}_3\text{R}$ currents (Fig. 4E, F). In this case, both Ca^{2+} responses and $5\text{HT}_3\text{R}$ currents were markedly reduced by ω -agatoxin IVA (Fig. 4E, F), consistent with the involvement of Ca_V channels in isovalerate-evoked EC cell Ca^{2+} responses (Fig. 2F). Importantly, none of these inhibitors directly affected $5\text{HT}_3\text{R}$ currents (Fig. S6C, D). Thus, Ca_V channels are required for serotonin release, likely mediating a local increase in intracellular Ca^{2+} near vesicular release sites, as observed for P/Q-type Ca_V channels in neuronal presynaptic terminals (Catterall et al., 2005b). Na_V channels are apparently not required for serotonin release, but may amplify responses to sub-threshold stimuli by generating action potentials.

EC cells regulate $5\text{HT}_3\text{R}$ -expressing nerves via synaptic-like contacts

$5\text{HT}_3\text{R}$ is robustly expressed by intrinsic neurons of the gut within submucosal and myenteric plexi, as well as by a subset of primary afferent sensory nerve fibers, including some that innervate the intestinal villi (Tecott et al., 1995). Moreover, pharmacological experiments with isolated (*ex vivo*) ileum strips suggest that AITC-evoked release of serotonin from EC cells produces intestinal contractions in a $5\text{HT}_3\text{R}$ -dependent manner (Nozawa et al., 2009). However, it is unclear whether such responses are mediated through diffuse, humoral spread of transmitter or a more direct, spatially restricted mechanism. In

fact, previous anatomical studies have suggested that peptide hormone-producing intestinal endocrine cells form synapses with neurons (Bohorquez et al., 2015). Using a reporter mouse expressing GFP under control of the 5HT₃R promoter (Vucurovic et al., 2010), we confirmed that many 5HT₃R-expressing nerve fibers innervate intestinal villi (Fig. 5A). Moreover, 5HT₃R-expressing fibers co-localized with synaptic markers, consistent with a neural origin (Fig. S7A). Interestingly, we observed multiple instances where 5HT₃R-expressing fibers appeared to make contact with the basolateral side of serotonin-expressing EC cells (Fig. 5A). Because EC cells express presynaptic P/Q-type Ca_v channels, are enriched for gene ontology categories related to neurotransmitter secretion, and express transcripts for several presynaptic markers (Fig. 5B, C), we wondered if this contact resembles a neural synapse. Indeed, EC cells demonstrated basolateral expression of the presynaptic marker, synapsin, and dense labeling of the postsynaptic marker, PSD-95, was observed immediately adjacent to EC cells (Fig. 5D). Thus, EC cells are in close proximity with 5-HT₃R-expressing nerve fibers and appear to form synaptic-like structures for transmitting signals in a restricted, point-to-point manner.

To determine if EC cells regulate activity of sensory fibers, we recorded from single, low threshold mechanosensitive pelvic fibers that project deep into the colonic tissue and access the mucosal layer containing EC cells. Epithelial application of norepinephrine evoked large responses in afferent nerves that were abolished by the TRPC4 inhibitor, ML204 (Fig. 6A). These responses were also blocked by the selective 5HT₃R antagonist, alosetron (Fig. 6A), indicating that they are mediated by 5HT₃R-expressing nerve fibers. Another EC cell agonist, isovalerate, evoked afferent nerve activity that was also blocked by alosetron (Fig. 6A). Neither norepinephrine nor isovalerate directly activated isolated, retrogradely-traced colonic sensory dorsal root ganglion (DRG) neurons, many of which were sensitive to the 5HT₃R-selective agonist, mCPBG (Fig. 6B). Thus, norepinephrine and isovalerate modulate 5HT₃R-expressing primary afferent nerve fibers via synaptically-coupled EC cells.

Chronic mechanical hypersensitivity contributes to the development of visceral pain syndromes, such as irritable bowel syndrome (Brierley and Linden, 2014). Interestingly, we found that epithelial application of norepinephrine or isovalerate markedly enhanced sensitivity of nerve fibers to mechanical stimulation of the colonic epithelium (Fig. 7 and Fig. S8). Norepinephrine modulatory effects were blocked by ML204 and responses to norepinephrine or isovalerate were abolished by alosetron (Fig. 7 and Fig. S8). These results are consistent with a role for EC cell-5HT₃R signaling in regulating mechanical sensitivity of the gut.

Discussion

EC cells as polymodal sensors of noxious stimuli

We show that EC cells are chemosensors that detect stimuli from three distinct sources including 1) ingested chemicals, 2) commensal organisms, and 3) endogenous regulatory pathways. Sensitivity to these agents is specified by receptors and transduction mechanisms that also contribute to other sensory or neural signaling systems. For example, TRPA1 is a well known somatosensory receptor for exogenous dietary irritants from mustard and allium plants, or endogenously produced inflammatory agents, such as 4-hydroxynonenal,

prostaglandins, and other lipid-derived metabolites (Bautista et al., 2006; Trevisani et al., 2007). TRPA1 has been implicated in visceral hypersensitivity, but this has for the most part been attributed to its function on colonic nerve fibers (Brierley et al., 2009). However, since the gut epithelium provides a barrier between the lumen and nerve fibers, EC cell-localized TRPA1 may serve as the primary detector of luminal irritants prior to direct sub-mucosal damage.

Effects of microbiota on the gastrointestinal system have been described at the organismal level, but underlying physiological mechanisms remain obscure (Schroeder and Backhed, 2016). For example, dietary fibers, proteins, and peptides are metabolized by commensal gut microbiota to produce volatile fatty acids, which in turn elicit diverse responses in the host through unknown mechanisms (Koh et al., 2016). Here we identify one such metabolite, isovalerate, as a potent EC cell stimulus that modulates sensory neurons via EC cell-neural signaling. Although isovalerate accounts for only a minor percentage of total fatty acid metabolites, high levels are toxic and associated with visceral pain and other gastrointestinal disorders, such as post-infectious irritable bowel syndrome (Brierley and Linden, 2014; Farup et al., 2016; Tanaka et al., 1966). EC cells may act as sensors for such potentially harmful dysbiosis.

As with other specialized sensory systems, EC cell chemosensation may adapt to detect stimuli that are most salient to an animal's physiologic or environmental conditions, such as specific diet, intestinal microbiota, etc. In this regard, it is interesting that mouse EC cells use Olf558 as a sensor for specific microbial metabolites, whereas mRNA encoding a different repertoire of olfactory receptors has been detected in human EC cells (Braun et al., 2007), possibly reflecting species-specific sensory tuning to suit distinct commensal relationships. An important and fascinating future goal is to explore signaling diversity and plasticity in EC cells from different species or under a variety of physiologic or pathophysiologic states.

EC cells are also sensitive to endogenous regulatory molecules, including stress-associated catecholamine neurotransmitters. Interestingly, norepinephrine is a potent bacterial stimulus that upregulates proliferation, virulence, and adherence of various pathogenic bacteria to influence the course of infection (Everest, 2007). Norepinephrine-mediated stimulation of EC cells may be protective by activating neural pathways that promote gastrointestinal motility to expel infectious microbes, metabolites, or harmful chemicals. However, EC cell-afferent nerve fiber stimulation resulting from prolonged infection or injury may also be maladaptive, eliciting chronic visceral hypersensitivity. Consistent with a link between gut catecholamine signaling and visceral pain, polymorphisms or deletions in *Adra2A* and *TRPC4* genes, respectively, are associated with visceral pain syndromes (Kim et al., 2004; Westlund et al., 2014). The gut is densely innervated by mechanosensory nerve fibers and enhanced afferent mechanical sensitivity is a hallmark of visceral pain (Brierley and Linden, 2014). We have shown that EC cell-neural chemosensory signaling cascades modulate mechanosensory function, establishing a direct mechanistic link between chemo- and mechanosensory elements in the gut. A recent study suggests that EC cells may themselves be mechanosensitive (Wang et al., 2017), but whether this contributes to mechanosensitivity of the gut is unknown.

EC cell signal transduction

Our results demonstrate that EC cells detect specific chemosensory stimuli using independent signaling pathways that converge on P/Q-type presynaptic voltage-gated Ca^{2+} channels to facilitate transmitter release onto afferent nerve fibers. Interestingly, while the Ca^{2+} -permeable transduction channels TRPA1 and TRPC4 each support large increases in intracellular Ca^{2+} , their activation is apparently insufficient to elicit serotonin release since inhibition of Ca_V channels completely blocks transmitter release while only slightly decreasing global Ca^{2+} responses. Voltage-gated sodium channels are not required for stimulus-evoked release, although they likely amplify responses by generating action potentials. These and other results suggest that Ca_V channels mediate obligatory changes in local Ca^{2+} , likely near the site of transmitter release, demonstrating the importance of cellular excitability and voltage-gated channels in EC cell function. Future studies of EC cell synaptic proteins, vesicular pools and docking mechanisms, and state-dependent plasticity will enhance our understanding of this process. Crosstalk among signal transduction cascades may also come into play. For example, we have shown that $\text{G}\alpha_i$ -coupled Adra receptors activate TRPC4, but $\text{G}\alpha_q$ -coupled receptors can also activate TRPC4, as well as TRPA1, providing opportunities for divergent and convergent effects of EC cell agonists on transmitter release.

Recent studies suggest a greater diversity of enteroendocrine cells than previously appreciated, arguing for the existence of genetically distinct EC cell subtypes (Diwakarla et al., 2017; Grun et al., 2015; Gunawardene et al., 2011). Indeed, we found that only a fraction of EC cells (7 of 62, Fig. 2A) were GABA-responsive, indicative of some degree of functional specification. However, other compounds elicited cellular responses in nearly all EC cells (Fig. 2A): AITC-evoked responses were observed in 15 of 15 high K^+ -responsive cells, norepinephrine responses in 30 of 30 cells, and isovalerate responses in 16 of 17 cells, suggesting that some functional attributes are conserved across most, if not all, putative EC cell subsets.

EC cell-neural communication

Signaling diversity or plasticity may also manifest at the level of neural circuitry. Our data show that EC cells make contact with $5\text{HT}_3\text{R}$ -expressing nerve fibers to mediate relatively local effects of serotonin. Consistent with previous reports (Aiken and Roth, 1992; Heitz et al., 1976), we found that EC cells are peptidergic, and that substance P expression is most prevalent in cells within crypts, decreasing upwards along the crypt-villus axis (Fig. S7B–D). Thus, EC cells may also release peptide transmitters to regulate synaptically-coupled sensory neurons. Moreover, hormone-producing enteroendocrine cells also form synaptic connections with nerve fibers (Bohorquez et al., 2015), although it is unknown if these neurons are of enteric or sensory origin. Therefore, sensory molecules could act on multiple enteroendocrine cell types to induce diverse signals (peptide hormones, serotonin) to regulate afferent nerve activity, conceivably transmitting specialized information through activation of specific fiber types or differential modulation of neural activity (Gribble and Reimann, 2016). Furthermore, discrete subclasses of DRG or vagal sensory nerves may communicate with specific resident EC cell populations (Williams et al., 2016). Finally, EC cell-derived serotonin may, in some cases, act on enteric neurons, immune cells, or be taken

up by circulating platelets to mediate diverse actions within or outside of the gut (Gershon, 2013; Veiga-Fernandes and Mucida, 2016).

EC cells and GI disorders

Alterations in EC cell-derived serotonin have been implicated in GI dysmotility, nausea, and visceral hypersensitivity disorders (Gershon, 2013; Mawe and Hoffman, 2013), and medications affecting norepinephrine or serotonin levels are associated with beneficial peripheral gut effects, such as relaxed fasting colonic muscle tone and reduction in mechanical hypersensitivity (Chial et al., 2003; Winston et al., 2010). Indeed, 5HT₃R-targeted therapeutics are used to treat chemotherapy-induced emesis, chronic nausea, and other visceral pain disorders (Mawe and Hoffman, 2013). Microbial metabolites also influence GI status and recent studies have noted beneficial effects of probiotics and differences in volatile fatty acids in patients with irritable bowel syndrome (Didari et al., 2015; Treem et al., 1996). Furthermore, intestinal inflammation is a key risk factor for the development of irritable bowel syndrome, and inflammatory molecules are associated with enhanced mechanical hypersensitivity of the gut (Brierley and Linden, 2014).

Our findings highlight EC cells as polymodal chemosensors that integrate extrinsic and intrinsic signals within the gut and convey this information to the nervous system. By exploiting intestinal organoid technology to access these rare, but important cells, we have gleaned new mechanistic insights into their function, which should facilitate EC cell-targeted therapeutics to treat irritable bowel syndrome and other disorders associated with gut hypersensitivity and pathophysiology.

Methods

Contact for reagent and resource sharing

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, David Julius (david.julius@ucsf.edu).

Experimental model and subject details

Animals—Mouse breeding, housing, and use was approved by the UCSF Animal Care and Use or University of Adelaide and Flinders University Animal Ethics Committees. Adult male mice of C57BL/6 background (Jackson Labs) aged 12–16 weeks with an average weight of ~29 grams were used for *ex vivo* afferent nerve recordings and colonic sensory neuron imaging. Male mice were used in all studies to account for effects from sex or genetic background. Animals were housed in groups (2–5 mice/cage) in a specific and opportunistic pathogen free facility, fed Jackson lab diet (5K52 JL RAT & MOUSE/AUTO 6F), provided with environmental enrichment (shelter, nesting material, etc.), and had normal immune status. Reporter mice were gifts from T. Schwartz (ChgA-GFP) and M. Scanziani (5HT₃R-GFP).

Intestinal organoids—Adult male ChgA-GFP mice aged 6–10 weeks were used to generate intestinal organoids, as previously reported (Sato et al., 2009). Briefly, the small intestine was isolated and washed with cold PBS and crypts were isolated following

dissociation in EDTA. Isolated crypts were suspended in Matrigel. Following polymerization, organoid growth media containing murine epidermal growth factor (PeproTech), noggin (PeproTech), and 10% R-spondin conditioned media was added and refreshed every 3–4 days. Organoids were maintained at 37°C, 5% CO₂ and propagated weekly. For Ca²⁺ imaging, Matrigel was removed from organoids, they were loaded with Fura-2AM, fenestrated by mechanical disturbance, and then immediately placed in the imaging chamber containing Cell-Tak (Corning)-coated coverslips. For electrophysiology, organoids were mechanically dissociated and placed on Cell-Tak coated coverslips in the recording chamber.

Cultured cells—Retrogradely traced colonic sensory neurons were isolated from adult male mice following injection of cholera toxin subunit B conjugated to AlexaFluor 488 (CTB-488; Invitrogen, Carlsbad, CA) at three sites sub-serosally within the wall of the distal colon. After 4 days, lumbosacral (LS) dorsal root ganglion neurons were isolated and cultured as previously described (Brierley et al., 2009). Briefly, mice were euthanized by CO₂ inhalation and lumbosacral dorsal root ganglia (DRGs) (L6-S1) from retrogradely traced mice were surgically removed and were digested with 4 mg/mL collagenase II (GIBCO, Life Technologies) plus 4 mg/mL dispase (GIBCO) for 30 minutes at 37°C, followed by 4 mg/mL collagenase II for 10 minutes at 37°C. Neurons were then mechanically dissociated into a single-cell suspension via trituration through fire-polished Pasteur pipettes. Neurons were resuspended in DMEM (GIBCO) containing 10% FCS (Invitrogen), 2mM L-glutamine (GIBCO), 100 μM MEM non-essential amino acids (GIBCO), 100 mg/ml penicillin/streptomycin (Invitrogen) and 100ng/ml NGF (Sigma). Neurons were spot-plated on coverslips coated with poly-D-lysine (800μg/ml) and laminin (20 μg/ml) and maintained at 37°C in 5% CO₂. Comparisons were made between retrogradely labeled colonic DRG neurons and non-labeled DRG neurons.

HEK293T (ATCC) were grown in DMEM, 10% fetal calf serum, and 1% penicillin/streptomycin at 37°C, 5% CO₂ and transfected using Lipofectamine 2000 (Invitrogen/Life Technologies) according to manufacturer's protocol. 1 μg of human Adra2A or TRPC4β was transfected for independent expression with 0.2 μg GFP. For coexpression experiments, equal concentrations of Adra2A and TRPC1, TRPC3, TRPC4, or TRPC6 were used. Constitutively active (Q205L) or dominant-negative (G203T) Gα_{i2} mutants were included in the transfection mix for some experiments. For Olfr558 experiments, 1.5 μg of Olfr558 was transfected with several constructs used enhance expression and signaling of olfactory receptors in heterologous systems: 1 μg of receptor-transporting protein 1 short (RTP1S), guanine nucleotide exchange factor B (Ric8b), and Gα_{olf15}, and 0.2 μg GFP for cellular identification. Mock transfection experiments were performed by transducing all constructs except Olfr558. Human Adra2A was from Genscript (Piscataway, NJ), Gα_{i2} mutants from cDNA Resource Center; TRPC1, TRPC3, and TRPC6 were cloned into pcDNA3 in the Julius lab; TRPC4 was a gift from J. Jeon and M. Zhu; Olfr558 constructs were gifts from H. Matsunami (Addgene) and were tagged with the first 20 residues of human rhodopsin to increase expression; RTP1S, Ric8b, and Gα_{olf15} were gifts from A. Chang. lentiCRISPR v2 was a gift from F. Zhang (Addgene).

Method details

CRISPR-mediated gene disruption—gRNA sequences were designed with the Cas9 design target tool (<http://crispr.mit.edu>) and inserted into the Cas9-containing lentiCRISPR v2 vector (Sanjana et al., 2014). Primers used to design the specific gRNA target were: Olfr558 forward (5' to 3') CACCGagcagctggcgcgtag; Olfr558 reverse (5' to 3') AAACctacggcatgccactgtgctC. Lentivirus was produced by transfecting HEK293T cells with psPAX2, pVSVG and LentiCRISPR v2 with Olfr558 gRNA using Fugene HD (Roche) according to the manufacturer's instructions. Virus was concentrated and re-suspended in organoid growth medium. As a control for both sequencing and functional experiments, organoids were infected with empty Cas9-containing LentiCRISPR v2 vector. Vector-infected organoids expressed wild-type Olfr558 sequence and exhibited similar isovalerate-induced Ca²⁺ responses compared with wild-type organoids, so were grouped with other controls in some analyses.

Two days before infection, intestinal organoids were grown in a 24-well culture plate with growth medium supplemented with 5 μM CHIR99021 (Sigma) and 10 mM nicotinamide (Sigma) to increase stem cell population. Stem cell-enriched organoids were broken down into single-cells, viral mix was added, and cells were transferred to a 48-well plate that was centrifuged at 600g for 60 min (spinoculation) and placed in the incubator for another 6 hr at 37 °C. Cells were collected, re-suspended in Matrigel, and transferred into a 24-well culture plate. After two days of recovery, selection was carried out using puromycin (6 μg/ml) for three days. After selection, growth medium was supplemented with 5 μM CHIR99021 and 10 mM nicotinamide. Organoids were then treated with TrypLE (Life Technologies) at 37 °C for 5 min to achieve single cells which were plated onto 96-well culture plates for clonal selection. Growth medium was supplemented with 5 μM CHIR99021, 10 μM Y-27632 (Sigma) and 10 mM nicotinamide during the first two days after plating to enrich for stem cells and prevent apoptosis. The medium was then changed to growth medium supplemented with 5 μM CHIR99021 and 10 mM nicotinamide for another three days and then normal growth medium afterwards. Single organoids were then collected and used for clonal expansion. To verify clonal populations genetic disruption, genomic DNA was isolated using QuickExtract DNA extraction solution (Epicenter) and PCR amplified using Phusion polymerase (NEB) and the following primers for Olfr558: F: cttgcatgcttctgacct; R: tgcagtggttctccattcca. Products were cloned into Topo vectors (Agilent) and sequenced.

Electrophysiology—Recordings were carried out at room temperature using a MultiClamp 700B amplifier (Axon Instruments) and digitized using a Digidata 1322A (Axon Instruments) interface and pClamp software (Axon Instruments). Data were filtered at 1 kHz and sampled at 10 kHz and leak-subtracted online using a P/4 protocol for voltage step protocols. Membrane potentials were corrected for liquid junction potentials. EC cells were identified by GFP expression and recordings were made using borosilicate glass pipettes polished to 7 – 9 MΩ. Recording pipettes used for HEK293 were 3 – 4MΩ. Unless stated otherwise, a standard Ringer's extracellular solution for EC and HEK293 cell experiments contained (mM): 140 NaCl, 5 KCl, 2 CaCl₂, 2 MgCl₂, 10 HEPES, 10 glucose, pH 7.4. Intracellular solution for recording K⁺ currents from EC cells contained: 140 K-gluc, 5 KCl, 1 MgCl₂, 10 K-EGTA, 10 HEPES, 10 sucrose, pH 7.2. Intracellular solution for

current-clamp recordings contained: 140 K-gluc, 5 NaCl, 1 MgCl₂, 0.02 K-EGTA, 10 HEPES, 10 sucrose, pH 7.2. Other EC cell and 5HT₃R recordings used the following intracellular solution: 140 CsMeSO₄, 5 NaCl, 1 MgCl₂, 10 Cs-EGTA, 10 HEPES, 10 sucrose, pH 7.2. Intracellular solution for Adra2A-expressing cells contained 0.1 Cs-EGTA. For EC recordings, holding potential was -90 mV and currents were elicited by 500ms ramps from -100 mV to +100 mV or 200ms steps in 10 mV increments. G-V relationships were derived from I-V curves by calculating $G = I_{Ca} / (V_m - E_{rev})$ and were then fit with a Boltzman equation. Voltage-dependent inactivation was measured during -10 mV voltage pulses following a series of 1 s prepulses ranging from -110 to 60 mV in 10 mV increments. Voltage-dependent inactivation was quantified as I / I_{max} , with I_{max} occurring at the voltage pulse following a -110 mV prepulse. Adra2A-associated experiments were carried out using a protocol that consisted of 10 s holding voltage at -60mV followed by a 500 ms ramp from -100 mV to +100 mV that returned to -60 for an additional 10s, and this protocol was repeated consecutively for ~10 min or more. For 5HT₃R biosensor recordings, whole-cell configuration was achieved and cells were lifted from coverslips and moved immediately adjacent to GFP-labeled EC cells. Voltage has held constant at -80 mV as solutions were washed on and off with local perfusion. Responses were normalized to peak current induced by mCPBG.

Calcium imaging—EC and HEK were loaded with 10 μM Fura-2-AM (Invitrogen) and 0.01% Pluronic F-127 (wt/vol, Invitrogen) for 1 h in Ringer's solution. 340 nm to 380 nm ratio was acquired using MetaFluor software. EC cells were identified by GFP expression and responses were normalized to increased fluorescence ratio elicited by high extracellular K⁺ (K⁺, 140 mM) at the end of the experiment. In most experiments, only one EC cell was identified in the field of view, thus we quantified data from single cells. In somewhat rare cases when two EC cells were observed in the same field of view, responses were averaged. Dorsal root ganglion neurons were cultured for 24 hours, incubated with 2.5 μM Fura2-AM and 0.02% (v/v) pluronic acid for 30 min at room temperature in modified Ringer's solution containing (mM): 145 NaCl, 5 KCl, 1.25 CaCl₂, 1 MgCl₂, 10 glucose, 10 HEPES. After a brief wash, coverslips were transferred to the recording chamber and Ca²⁺ responses were measured at room temperature. Colonic DRG neurons were identified by the presence of the 488 tracer and viability was verified by responses to 25 mM KCl. All pharmacological agents were delivered by local perfusion with exception of 1 μM U73122, 100 μM gallein, 200 ng/ml cholera toxin, or 200 ng/ml pertussis toxin, 10 μM SQ22536, which were preincubated. Associated vehicle control experiments were performed. In experiments using HEK293, construct-expressing cells identified by GFP expression were quantified and responses were normalized to maximal responses elicited by 1 μM ionomycin at the end of the experiment.

Concentrations and abbreviations of molecules used in Ca²⁺ imaging screening (in μM): 1 Capsaicin, 500 allyl isothiocyanate (AITC), 50 1-(m-chlorophenyl)-biguanide (mCPBG), 1 icilin, 200 N-butyryl-L-Homoserine lactone (C4-HSL), 200 N-hexanoyl-L-Homoserine lactone (C6-HSL), 200 N-3-oxododecanoyl-L-Homoserine lactone (3OC12-HSL), 1 or 10 N-Formylmethionine-leucyl-phenylalanine (fMFL), 50 μg/ml lipopolysaccharide (LPS) from E. Coli, 500 indole, 500 sodium propionate, 500 sodium acetate, 500 sodium butyrate,

500 isobutyrate, 500 isovalerate, 500 sodium deoxycholate, 1 substance P, 100 histamine, 1000 glutamate, 100 tryptamine, 100 serotonin, 100 glycine, 100 gamma-aminobutyric acid (GABA), 100 dopamine, 100 epinephrine, 100 norepinephrine. Unless stated otherwise, concentrations of other pharmacological agents (in μM): 0.5 tetrodotoxin (TTX), 10 nifedipine, 0.3 ω -agatoxin IVA, 0.3 ω -conotoxin, 5 mibefradil, 1 epinephrine, 1 norepinephrine, 5 yohimbine, 10 isoproterenol, 5 prazosin, 5 clonidine, 5 propranolol, 10 phenylephrine, 5 U73122, 100 gallein, 200ng/ml pertussis toxin (PTX), 200ng/ml cholera toxin (CTX), 50 2-aminoethoxydiphenyl borate (2-APB), 10 ML204. Most drugs were from Tocris, HSLs and 4-hydroxynonenal were from Cayman Chemical, volatile fatty acids were from Sigma.

Transcriptome sequencing and analysis—Intestinal epithelial cells from organoids were dissociated and immediately sorted by fluorescence-activated cell sorting (FACS) by the Laboratory for Cell Analysis at UCSF. ~1% of total epithelial cells were GFP⁺ and collected. The remaining GFP⁻ cells were retained for comparison. RNA from GFP⁺ and GFP⁻ subgroups was then extracted and prepared for cDNA library generations using the SMARTer Ultra Low Input RNA kit followed by the Low Input Library Prep Kit (version 2, Clontech Laboratories, Inc.). cDNA quality was assessed via bioanalyzer using the High Sensitivity DNA kit (Agilent Technologies), and high quality samples were preserved for sequencing.

PolyA cDNA libraries were sequenced on the Illumina Hi-Seq 4000 platform (QB3 Vincent J. Coates Genomic Sequencing Library), generating 150 bp paired-end reads. More than 100M reads were obtained. The quality of raw sequence reads was analyzed via FASTQC. Adapters were trimmed using Scythe, and sequence read ends were trimmed using Sickle. Reads were then aligned to the annotated mouse reference genome (mm10) using TopHat2 (version 0.7). Transcripts were assembled and relative abundance was estimated using Cufflinks and Cuffdiff tools. Gene ontology-based (GO) analyses were carried out using DAVID (version 6.8) to categorize the top ~1000 transcripts annotated with ENSEMBL gene IDs that showed the greatest fold change between GFP⁺ and GFP⁻ samples. The “biological process” set of GO terms was used in functional annotation of the enriched transcripts in the GFP⁺ sample over the GFP⁻ sample, which was set as the background.

Histology—Immunofluorescence (IF) was performed using 5 μm cryosections. Blocking was performed with 10% normal serum corresponding to secondary antibody species in 0.1% Triton-X and PBS at room temperature for 60 minutes. Primary antibodies were incubated overnight at 4 °C at the indicated dilutions. Antibodies used were against ChgA (1:200, Santa Cruz), serotonin (1:10,000, Immunostar), Adra2A (1:200, Affinity Bioreagents), tyrosine hydroxylase (1:500, Millipore), Synapsin (1:500, from R. Edwards), PSD-95 (1:200, Neuromab), Lysozyme (1:200, Dako), GLP-1 (1:200, Abcam), Substance P (1:1000, Penninsula). Alexa Fluor-conjugated secondary antibodies were used at 1:300-1000 (Millipore). *In situ* hybridization histochemistry was performed using digoxigenin- and fluorescein-labelled cRNA for mouse TRPC4 or Olfr558. Probes were generated by T7/T3 *in vitro* transcription reactions using a 500-nucleotide fragment of TRPC4 (nucleotides 1553 to 2053), and a 500-nucleotide fragment of Olfr558 cDNA (nucleotides 1000 to 1500).

Hybridization was developed using anti-digoxigenin and anti-fluorescein Fab fragments, followed by incubation with FastRed and streptavidin-conjugated Dylight 488 according to published methods (Ishii et al., 2004). Epifluorescence imaging was performed on an Olympus IX51 microscope equipped with a DP71 imager and Nikon Eclipse Ti with a DS-Qi2 imager. Confocal imaging was performed on Nikon Ti microscope with Yokogawa CSU-22 spinning disk. Images were assembled in Photoshop and ImageJ. Surface rendering was performed using UCSF Chimera.

Single nerve fiber recordings of pelvic colonic mucosal afferents—C57BL/6J male mice were humanely euthanized by CO₂ inhalation. The colon and rectum with attached pelvic nerves were removed and recordings from mucosa afferents were performed as previously described (Brierley et al., 2004). Briefly, the colon was removed and pinned flat, mucosal side up, in a specialized organ bath. The colonic compartment was superfused with a modified Krebs solution (in mM: 117.9 NaCl, 4.7 KCl, 25 NaHCO₃, 1.3 NaH₂PO₄, 1.2 MgSO₄, 2.5 CaCl₂, 11.1 D-glucose), bubbled with carbogen (95% O₂, 5% CO₂) at a temperature of 34°C. All preparations contained the L-type calcium channel antagonist nifedipine (1 μM) to suppress smooth muscle activity and the prostaglandin synthesis inhibitor indomethacin (3 μM) to suppress potential inhibitory actions of endogenous prostaglandins. The pelvic nerve bundle was extended into a paraffin-filled recording compartment in which finely dissected strands were laid onto a mirror, and a single fiber placed on the platinum recording electrode. Action potentials recorded in response to mechanical or chemical stimuli were discriminated as single units based on Waveform, amplitude and duration using spike software (Cambridge Electronic Design, Cambridge, UK).

Colonic afferents were classified by identifying receptive fields by systematically stroking the mucosal surface with a still brush to activate all subtypes of mechanoreceptors. Categorization of afferents properties was in accordance with our previously published classification system (Brierley et al., 2004). In short, pelvic mucosal afferents respond to fine mucosal stroking (10 mg von Frey hairs; vfh), but not to circular stretch. Stimulus–response functions were constructed by assessing the total number of action potentials generated in response to mechanical stimuli (mucosal stroking with 10, 200, 500 and 1000 mg vfh). Norepinephrine (NE, 1 μM) was applied for 15 minutes via a small metal ring placed over the receptive field of interest and the TRPC4 inhibitor ML204 (10 μM) or the 5-HT₃R antagonist alosetron (10 μM) were pre-incubated for 10 minutes prior, and co-applied with NE. This route of administration has been previously shown to reproducibly activate afferent fibers (Brierley et al., 2009).

Quantification and statistical analysis

Data were analyzed with Clampfit (Axon Instruments) or Prism (Graphpad) and are represented as mean ± sem and n represents the number of cells or independent experiments. Data were considered significant if $p < 0.05$ using paired or unpaired two-tailed Student's *t*-tests or one- or two-way ANOVAs. Statistical parameters are described in figure legends. All significance tests were justified considering the experimental design and we assumed normal distribution and variance, as is common for similar experiments. Sample sizes were chosen

based on the number of independent experiments required for statistical significance and technical feasibility.

Data availability

Deep sequencing data have been deposited in Gene Expression Omnibus (GEO) database repository with accession number GSE98794. All other data are available from the authors upon request.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Highlights

- Enterochromaffin (EC) cells are excitable and express voltage-gated ion channels
- EC cells use sensory receptors to detect irritants, metabolites, and catecholamines
- EC cell activation leads to voltage-gated Ca^{2+} channel-dependent serotonin release
- EC cells modulate sensory nerves via serotonin receptors and synaptic connections

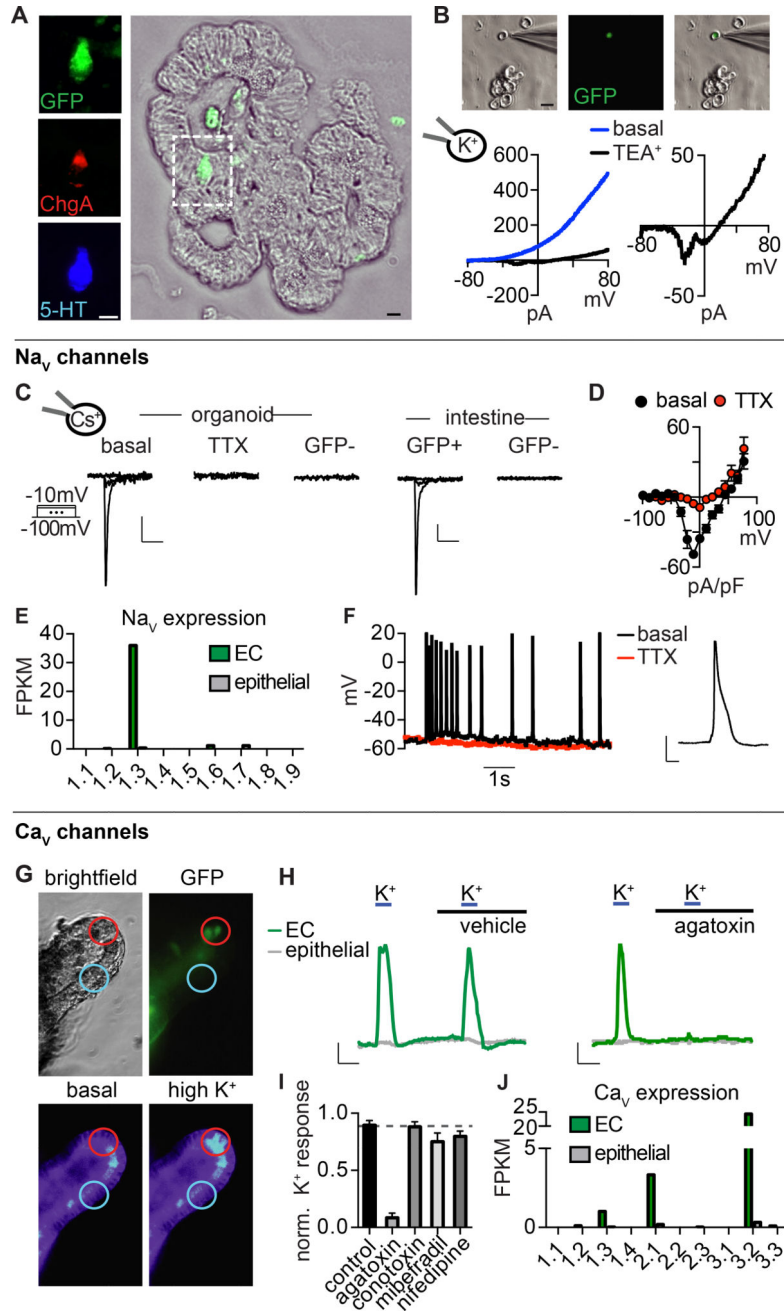


Figure 1. Enterochromaffin cells are electrically excitable

A. Co-localization of chromogranin A-driven GFP reporter (ChgA-GFP, green), ChgA (red), and serotonin (5-HT, blue) labels enterochromaffin (EC) cells in intestinal organoids. Scale bar: 10 μ m.

B. Dissociated EC cell (green) in a representative patch-clamp experiment. Scale bar: 10 μ m. In response to a voltage ramp, the representative K⁺ current was blocked by 10 mM TEA⁺ to reveal a voltage-activated inward current. Representative of n=4 cells.

C. Voltage-gated currents in EC cells were inhibited by the Na_v antagonist tetrodotoxin (TTX, 500nM). Scale bars: 50pA vertical, 10ms horizontal.

- D.** Average current-voltage relationship. $n=7$. $p<0.0001$ for basal versus TTX. Two-way ANOVA with post-hoc Bonferroni test.
- E.** mRNA expression profile of Na_v pore-forming subunits in EC cells (green) compared with other intestinal epithelial cells (grey). Bars represent fragments per kilobase of exon per million fragments mapped (FPKM).
- F.** Spontaneous action potentials measured at resting membrane potential were inhibited by TTX. Representative of $n=4$. *Inset*: representative action potential, scale bar: 20mV, 10ms.
- G.** Representative calcium (Ca^{2+}) imaging experiment from EC cells (GFP, green) in an intestinal organoid. High extracellular K^+ increased cytosolic Ca^{2+} , indicated by a change in fluorescence ratio of Fura-2AM.
- H.** Ca^{2+} responses to K^+ -elicited depolarization in EC (green) or neighboring cells (grey). The P/Q-type Ca_v inhibitor ω -agatoxin IVA abolished responses. Scale bar: 0.25 Fura-2 ratio, 50s.
- I.** Pharmacological profile of Ca_v -mediated responses. $n=6$ per condition. Data represented as mean \pm sem. $p<0.0001$ for control versus 300nM ω -agatoxin IVA. One-way ANOVA with post-hoc Bonferroni test. All data represented as mean \pm sem.
- J.** mRNA expression profile of Ca_v pore-forming subunits in EC cells (green) compared with other intestinal epithelial cells (grey).

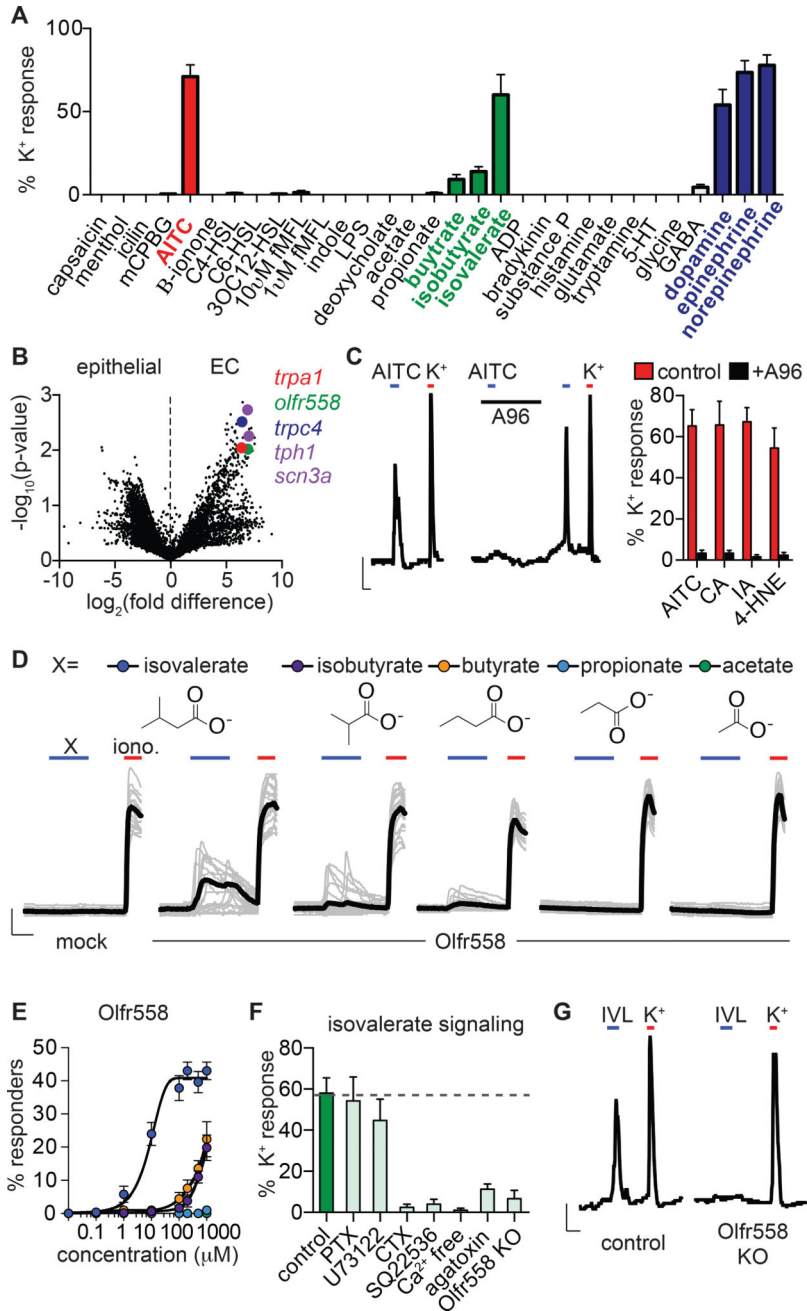


Figure 2. Enterochromaffin cells use TRPA1 as an irritant receptor and Olfr558 as a metabolite sensor

A. Sensory molecule screen for EC cell-specific Ca²⁺ responses. High K⁺ was added at the end of each experiment to induce maximal Ca²⁺ responses used for normalization. n=6–62 per condition. Data represented as mean ± sem.

B. mRNA expression profile of EC cells compared with other intestinal epithelial cells shown as a volcano plot. *Trpa1* (red), *olfr558* (green), and *trpc4* (blue) were among the most enriched transcripts that encode sensory receptors or channels. EC cell marker *tph1* and Nav1.3 pore-forming subunit *scn3a* are shown for comparison (purple).

C. AITC (150 μ M)-elicited Ca²⁺ responses were inhibited by the TRPA1 antagonist A967079 (A96, 10 μ M). Scale bar: 0.1 Fura-2 ratio, 50s. Average peak Ca²⁺ responses evoked by TRPA1 agonists AITC, cinnamaldehyde (CA, (150 μ M), iodoacetamide (IA, 150 μ M), or 4-hydroxynonenal (4-HNE, 200 μ M) were inhibited by A96 (10 μ M). n=5 per condition. p<0.0001 for agonists versus agonists + A96, two-way ANOVA with post-hoc Bonferroni test.

D. Ca²⁺ responses elicited by metabolites (200 μ M) in HEK293 cells expressing Olfr558. Ionomycin (iono, 1 μ M) was added at the end of each experiment to induce maximal Ca²⁺ responses. Black traces represent an average of all cells in the field shown in grey. Scale bar: 0.2 Fura-2 ratio, 50s.

E. Dose-response comparing isovalerate (blue), isobutyrate (purple), butyrate (orange), propionate (light blue), or acetate (green) represented as % of cells that responded to the indicated concentration of each compound. n=6 per condition. EC50 for isovalerate was 8.92 μ M with a 95% confidence interval of 7.32 to 10.51 μ M.

F. Isovalerate-evoked Ca²⁺ responses in EC cells. n=5 per condition. p<0.001 for control (vehicle-treated or empty Cas9-containing vector-infected organoids) versus cholera toxin (CTX), adenylyl cyclase inhibitor SQ22536 (10 μ M), Ca²⁺ free extracellular solution, ω -agatoxin IVA (300nM), Olfr558 knockout (KO). All data represented as mean \pm sem. n=5–8 per condition, one-way ANOVA with post-hoc Bonferroni test.

G. Isovalerate (IVL, 200 μ M)-evoked responses were absent in Olfr558 knockout (KO) ChgA-GFP organoids generated using CRISPR. Scale bar: 0.1 Fura-2 ratio, 50s.

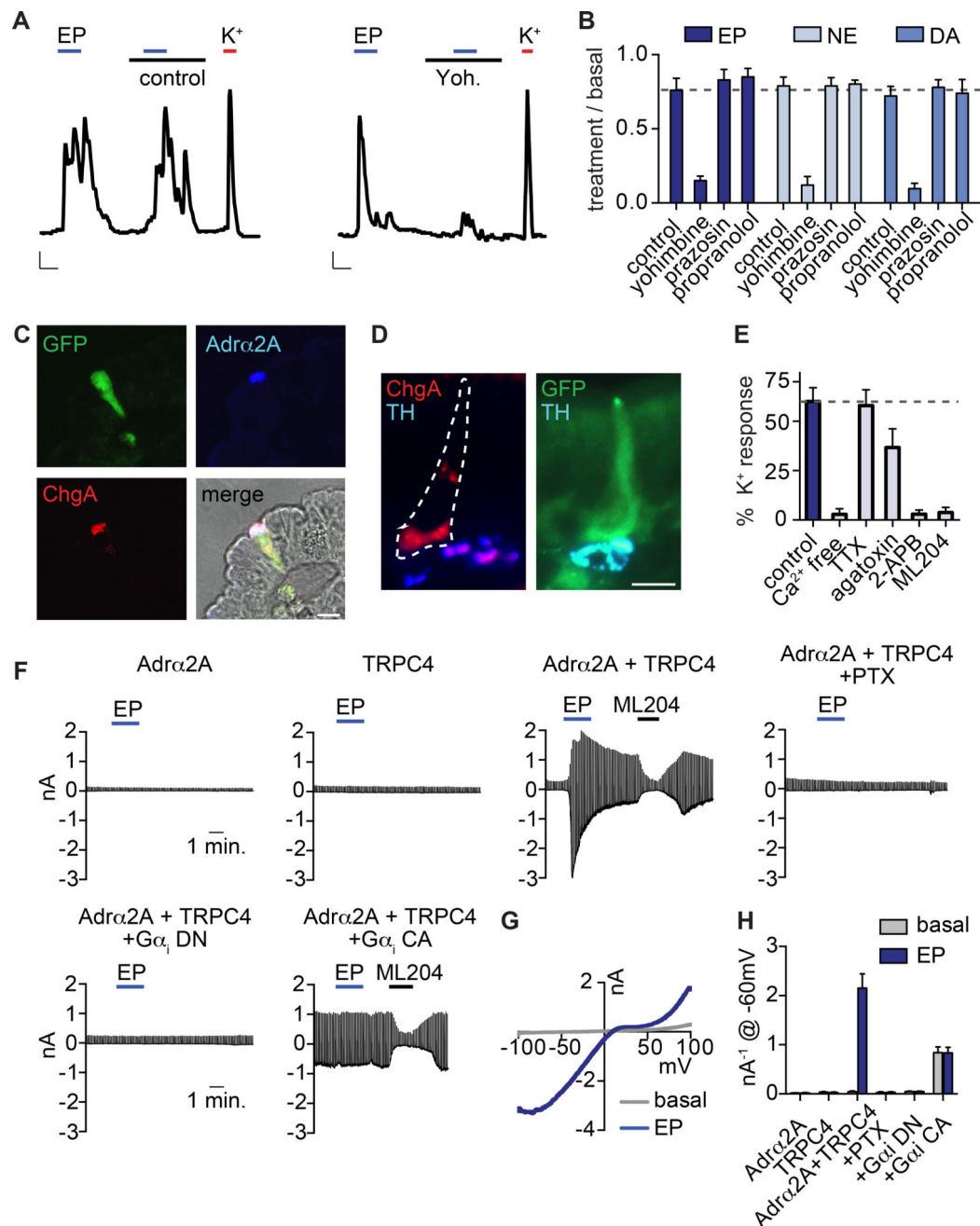


Figure 3. Adra2A and TRPC4 form a catecholamine-sensitive signaling cascade in enterochromaffin cells

A. Epinephrine (EP, 1 μ M)-evoked Ca²⁺ responses were blocked by the adrenoreceptor α 2 (Adra2) antagonist yohimbine (yoh, 5 μ M). Scale bars: 0.1 Fura-2 ratio, 50s.

B. Average peak catecholamine responses were inhibited by the Adra2 antagonist yohimbine, but not the Adra1 antagonist prazosin (5 μ M) or the Adr β antagonist propranolol (5 μ M). n=5 per condition. p<0.0001 for control versus yohimbine for EP (1 μ M), norepinephrine (NE, 1 μ M), dopamine (DA, 100 μ M). Two-way ANOVA with post-hoc Bonferroni test.

C. Adra2A (blue) localized to the basolateral side of EC cells (indicated by ChgA in red or GFP reporter) and was specific among intestinal epithelial cells. Scale bar: 10 μ m.

D. Tyrosine hydroxylase (TH, blue), a marker for norepinephrine-producing sympathetic nerve fibers, localized on the basolateral side of EC cells (indicated by ChgA in red or GFP reporter). Scale bar: 10 μ m.

E. Pharmacological profile of EP responses. n=7 per condition. p<0.0001 for control versus Ca²⁺ free, TRPC inhibitor 2-APB (50 μ M), TRPC4 inhibitor ML204 (10 μ M); p<0.05 for control versus ω -agatoxin IVA (300nM). One-way ANOVA with post-hoc Bonferroni test.

F. EP-elicited currents were elicited from HEK293 coexpressing Adra2A and TRPC4, but not cells independently expressing Adra2A or TRPC4. EP-elicited currents were inhibited by pertussis toxin (PTX, 200ng/ml) or coexpression of dominant-negative (DN) G α_i . Coexpression of constitutively-active (CA) G α_i induced ML204-sensitive activity that occluded EP-elicited currents.

G. Representative current-voltage relationship shows the peak EP response (blue) and basal current (grey) from the representative cell expressing Adra2A and TRPC4 shown in F.

H. Average peak current amplitude recorded at -60mV before (basal, grey) or during EP (blue) application. n=6 per condition. All data represented as mean \pm sem. p<0.0001 for basal versus epinephrine-evoked currents in Adra2A and TRPC4, two-way ANOVA with post-hoc Bonferroni test.

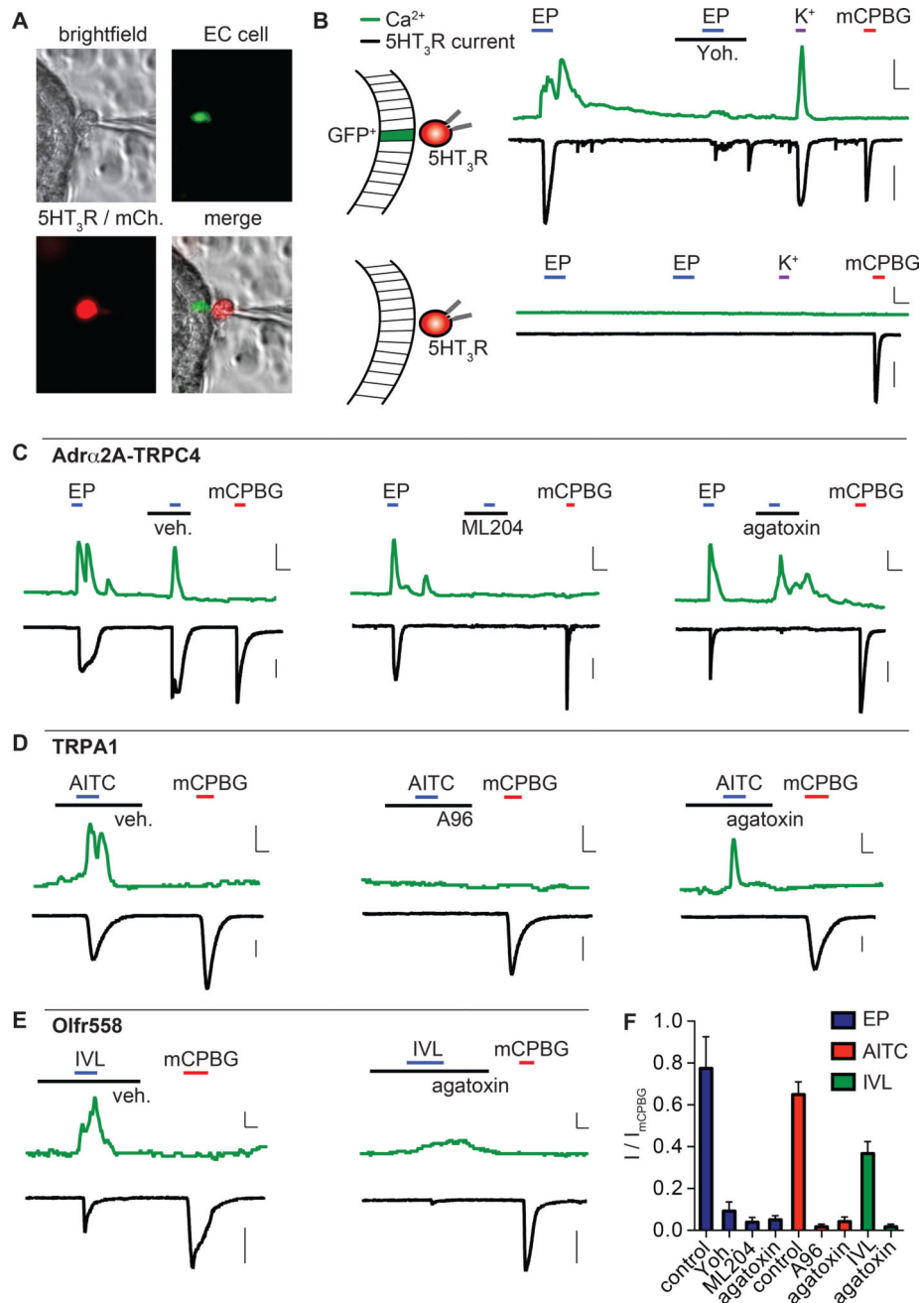


Figure 4. Enterochromaffin cell activation mediates Ca_V-dependent 5-HT release

A. Representative 5-HT “biosensor” experiment. 5HT₃R-expressing HEK293 (mCherry, red) adjacent to an EC cell (GFP, green) for simultaneous Ca²⁺ measurements from EC cells and whole-cell current measurements from biosensor cells.

B. Epinephrine (EP, 1 μ M) or high extracellular K⁺ induced a Ca²⁺ response in EC cells that correlated with a large 5HT₃R current in biosensor cells. EP responses were inhibited by yohimbine (yoh, 5 μ M). The 5HT₃R agonist mCPBG (10 μ M) elicited a large biosensor current, but no EC cell Ca²⁺ response. When biosensor cells were moved away from EC cells, neither epinephrine nor K⁺ induced Ca²⁺ responses in GFP⁻ epithelial cells or

biosensor currents, but mCPBG elicited a large 5HT₃R current. Scale bars: 0.6 Fura-2 ratio, 50s, 500pA.

C. EP-evoked Ca²⁺ responses and 5HT₃R currents were not affected by vehicle but were blocked by the TRPC4 inhibitor ML204 (10μM). The Ca_V inhibitor ω-agatoxin IVA (300nM) slightly reduced Ca²⁺ responses and abolished 5HT₃R currents. Scale bars: 0.3 Fura-2 ratio, 50s, 500pA.

D. AITC (150μM)-evoked Ca²⁺ responses and 5HT₃R currents were blocked by the TRPA1 antagonist A967079 (A96, 10μM). ω-agatoxin IVA (300nM) did not significantly affect Ca²⁺ responses, but abolished 5HT₃R currents. Scale bars: 0.1 Fura-2 ratio, 25s, 500pA.

E. Isovalerate (IVL, 200μM)-evoked Ca²⁺ responses and 5HT₃R currents. ω-agatoxin IVA (300nM) inhibited Ca²⁺ responses and abolished 5HT₃R currents. Scale bars: 0.1 Fura-2 ratio, 25s, 500pA.

F. Average agonist-evoked biosensor currents normalized to mCPBG-induced current (I_{mCPBG}). n=4 – 5 per condition. Data represented as mean ± sem. Responses to epinephrine (EP, blue), AITC (red), and isovalerate (IVL, green) in the presence of indicated antagonists. p<0.001 for control versus treatments. One-way ANOVA with post-hoc Tukey's test.

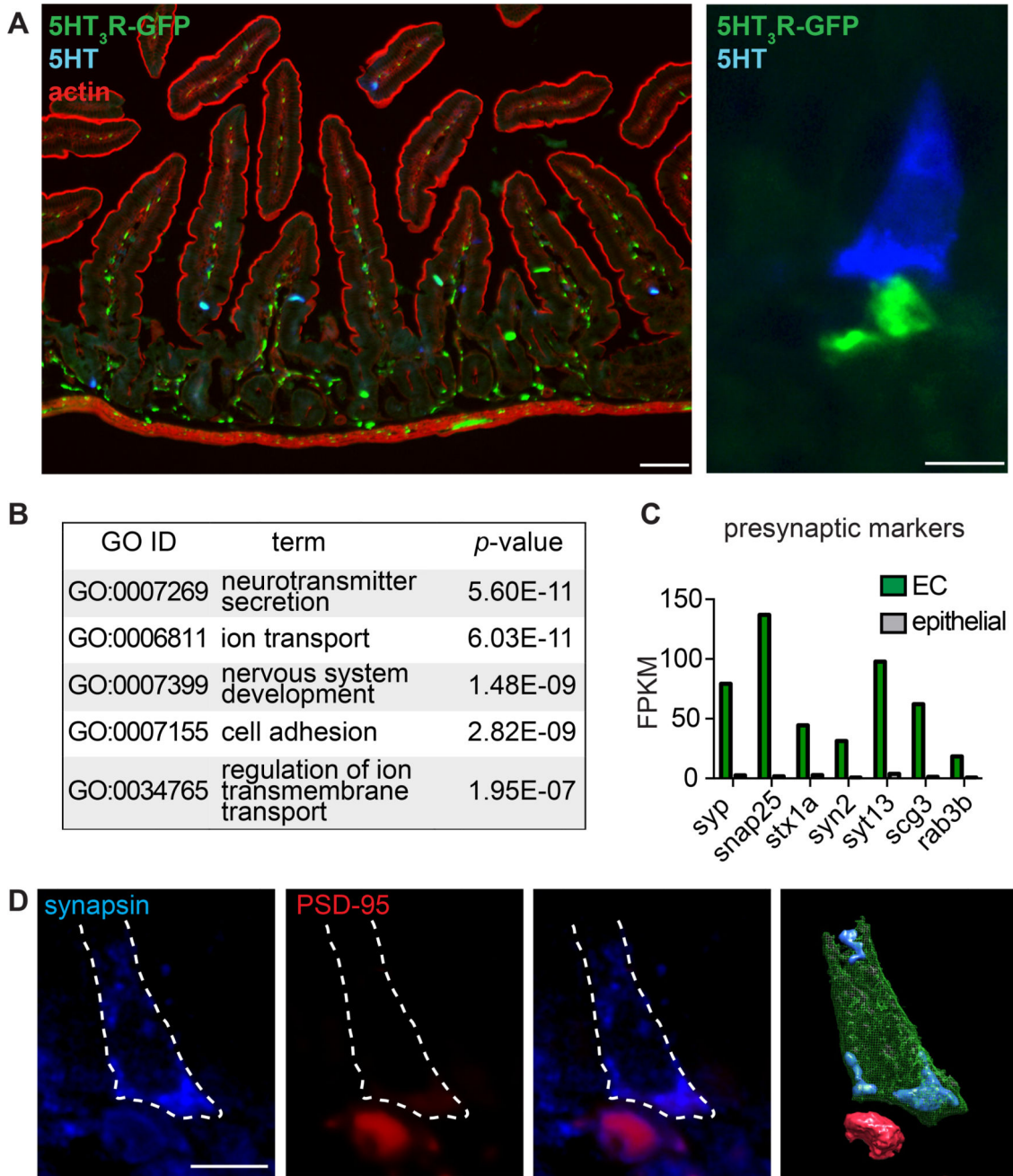


Figure 5. Enterochromaffin cells form synaptic-like contacts with 5HT₃R-expressing nerve fibers
A. (*Left*) Representative jejunal cryosection showing 5HT₃R-expressing fibers (green) innervating intestinal villi near serotonin-expressing EC cells (5-HT, blue) with actin staining (red) to demonstrate intestinal architecture. Scale bar: 50 μ m. (*Right*) Representative image demonstrating proximity between a 5-HT-positive EC cell (blue) and 5HT₃R-expressing fiber (green, basolateral side). Scale bar: 10 μ m.
B. Top 5 enriched Gene ontology (GO) categories in EC cells compared with other intestinal epithelial cells.

C. Presynaptic marker mRNA expression profile in EC cells (green) versus other intestinal epithelial cells (grey). Bars represent fragments per kilobase of exon per million fragments mapped (FPKM).

D. A 5 μm section of intestinal epithelium showing a representative EC cell that expressed the presynaptic marker synapsin (blue, basolateral side) and made contact with a postsynaptic marker-positive fiber (PSD-95, red). Cell body is outlined (dashed white line). Three-dimensional rendering of EC cell (green) with synapsin-positive vesicles (blue) near postsynaptic-like structure (red). Scale bar: 10 μm .

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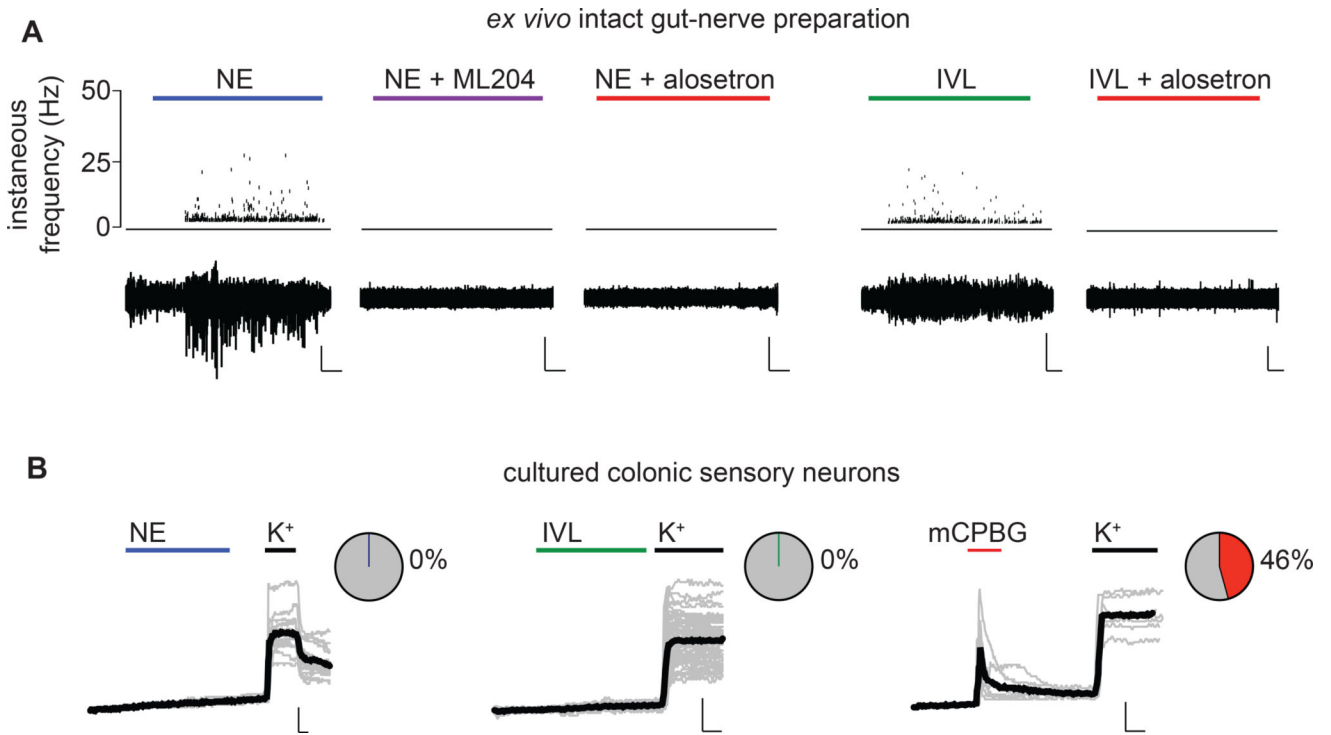


Figure 6. Enterochromaffin cells modulate 5HT₃R-expressing afferent nerves

A. Representative recordings from single mucosal afferent nerve fibers innervating intact colonic epithelium in an *ex vivo* preparation. Norepinephrine (NE, 1 μ M) applied to the epithelium elicited chemosensory responses that were blocked by the TRPC4 inhibitor ML204 (10 μ M) or the 5HT₃R antagonist alosetron (10 μ M). Isovalerate (IVL, 200 μ M) also evoked alosetron-sensitive afferent activity. Scale bars: 500 μ V, 50s. Representative of n=8–9 per condition. $p < 0.0001$ for number of action potentials measured in response to NE (321.5 \pm 55.4 spikes, 4/8 responsive fibers) versus NE+ML204 (0 spikes, 0/8 responsive fibers) or NE+alosepron (0 spikes, 0/9 responsive fibers), one-way ANOVA with post-hoc Bonferroni test. $p < 0.01$ for number of action potentials measured in response to IVL (648.7 \pm 339.3 spikes, 3/8 responsive fibers) versus IVL+alosepron (0 spikes, 0/8 responsive fibers).

B. The 5HT₃R agonist mCPBG (10 μ M), but not NE (1 μ M) or isovalerate (IVL, 200 μ M), evoked representative Ca²⁺ responses in retrogradely-labeled colonic sensory neurons isolated from lumbosacral dorsal root ganglia. All neurons quantified responded to high extracellular K⁺. Black traces represent an average of all cells in the field shown in grey. Scale bar: 0.1 Fura-2 ratio, 60s. Responsive neurons: n=0/16 for NE, n=0/62 for IVL, n=16/35 for mCPBG.

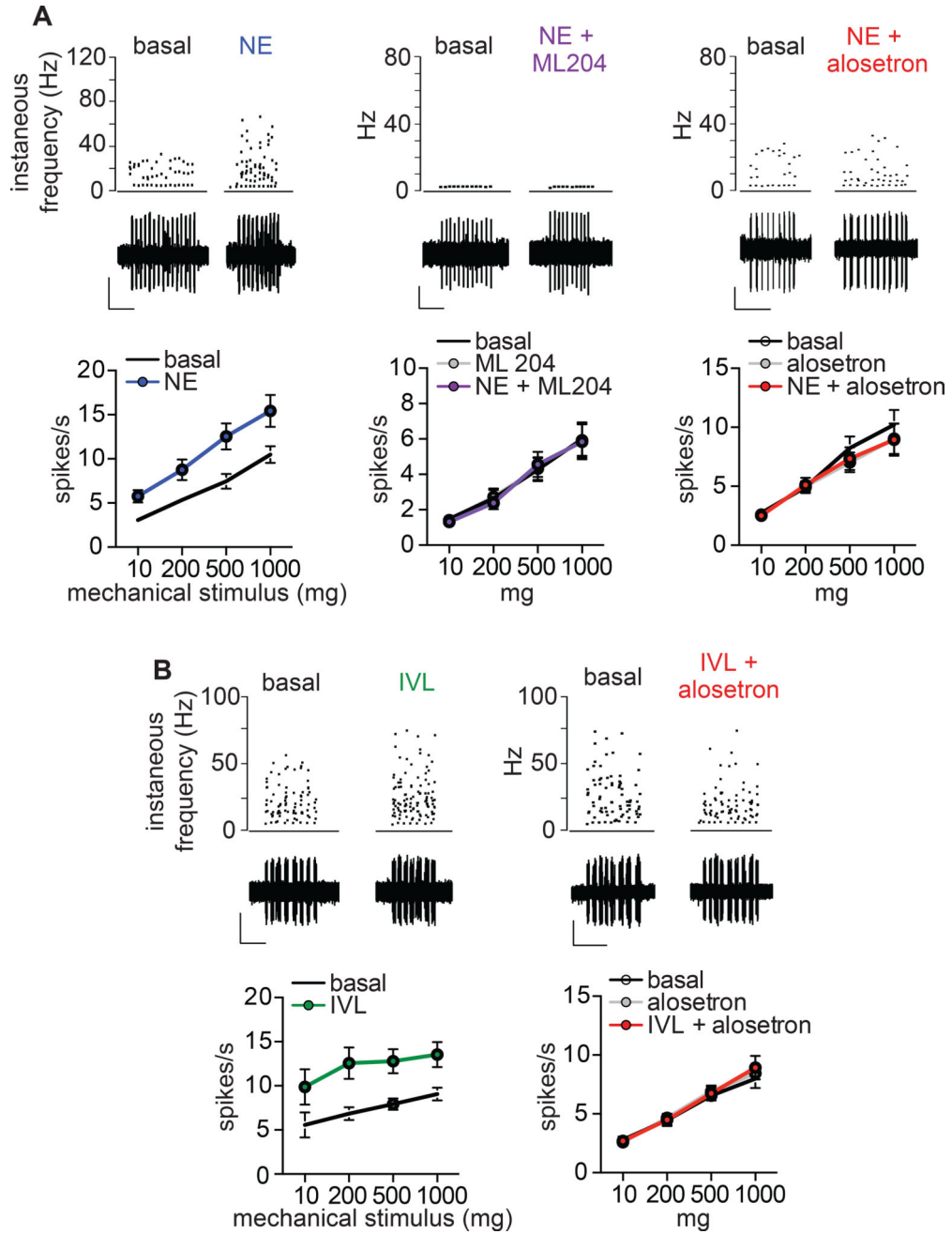


Figure 7. Enterochromaffin cells induce mechanical hypersensitivity of colonic afferents
A. Representative mechanical responses from single low-threshold mechanoreceptive mucosal afferent fibers elicited by a 10mg von Frey hair stimulus to epithelium. Mechanical responses were enhanced following epithelial treatment with norepinephrine (NE, 1 μ M) and hypersensitivity was blocked by the TRPC4 inhibitor ML204 (10 μ M) or 5HT₃R antagonist alosetron (10 μ M). n=8–9 per condition. Scale bars: 400 μ V, 10s. $p < 0.0001$ for contribution of treatment to series variance for NE versus basal and no significant difference with NE +ML204 or alosetron, two-way ANOVA with post-hoc Bonferroni test.

B. Afferent mechanosensory responses were enhanced following epithelial treatment with isovalerate (IVL, 200 μ M) and hypersensitivity was blocked by alosetron (10 μ M). A 500mg von Frey hair was used as an epithelial mechanical stimulus for representative traces. n=8–9 per condition. Scale bars: 500 μ V, 10s. $p < 0.0001$ for contribution of treatment to series variance for IVL versus basal and no significant difference with IVL+alosectron, two-way ANOVA with post-hoc Bonferroni test. All data represented as mean \pm sem.

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