Uroguanylin: how the gut got another satiety hormone

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Prouroguanylin is a gastrointestinal paracrine signal and prohormone that is secreted after nutrient ingestion. In this issue of the JCI, Valentino et al. show that prouroguanylin is converted to uroguanylin in the CNS, which can activate guanylyl cyclase 2C (GUCY2C) receptors in the brain to reduce food intake in mice. This 16–amino acid residue peptide is a novel component of the gut-brain axis that represents a new and unique opportunity to manipulate gut-brain signaling for therapeutic intervention in obesity.

Life as an endocrinologist is getting complicated these days. A few decades ago, there were just a small number of endocrine organs, including the thyroid, adrenals, and pancreas. Over the past 20 years, the list of organs that are considered major endocrine organs has expanded to include adipose tissue, liver, bone, and muscle, among others. The endocrine productivity of these organs, however, cannot compete with that of the gastrointestinal (GI) tract. From the stomach to the rectum, there are numerous distinct cell populations that secrete a wide range of hormones involved in a diverse array of functions (1).

One of the key functions of GI hormones is the regulation of food intake and the distribution of ingested nutrients (2). Particularly intriguing data highlighting the important role of GI hormones in this context come from the assessment of patients undergoing bariatric surgery, for whom a variety of manipulations of the GI tract result in profound weight loss and the resolution of comorbid conditions, such as type 2 diabetes (3). While it remains a possibility that the changes that result from bariatric surgery are the direct result of the physical restriction to food intake and nutrient absorption and/or malabsorption that may accompany such procedures, it has become increasingly accepted that modulation of the levels of a wide range of factors secreted by the GI tract may have a crucial role in the effectiveness of these therapies (4).

In the clinical arena, GI hormones have a potential role as the molecular signals that mediate the metabolic benefits of bariatric surgeries, and several new GI hormone–based drugs are now available to the patient or are in clinical testing for the treatment of metabolic diseases. A key role for the GI tract, as both potential cause and cure for escalating levels of metabolic diseases therefore seems likely, making a thorough understanding of its complex endocrine functions critical for our success in the fight against obesity and type 2 diabetes. In this issue of the JCI, Valentino et al. report a new player involved in the cornucopia of hormones that is the GI endocrine system (5). Identifying this novel system opens up new possibilities for GI-based therapies for helping obese individuals lose weight.

Guanylin and uroguanylin: new gut hormones

Guanylin and uroguanylin are peptides of 15– and 16–amino acids in length, respectively. They are both secreted by intestinal epithelial cells as prohormones, which require enzymatic conversion into active hormones. To date, their function has been thought to be largely paracrine. Upon secretion into the lumen of the GI tract, they act on guanylyl cyclase 2C (GUCY2C) receptors on intestinal epithelial cells, increasing intracellular cyclic GMP (cGMP). Interestingly, GUCY2C receptors also are the target for heat-stable enterotoxins and upon activation can decrease intestinal fluid absorption, which can lead to diarrhea (6, 7). cGMP, however, has also been implicated in the regulation of nutrient intake in invertebrate models (8). Therefore, Valentino and colleagues hypothesized that uroguanylin or guanylin might act as part of the gut-brain axis that regulates ingestion, energy homeostasis, and body weight (5). Consistent with this hypothesis, they found that mice engineered to globally lack the GUCY2C receptor carried more body fat as a direct result of increased food intake. These observations were further corroborated when they observed that systemic administration of a GUCY2C agonist reduced food intake in wild-type mice but not GUCY2C-deficient mice.

Prouroguanylin: a postprandial hormone that is activated in the brain

One of the key questions after having determined that signaling via the GUCY2C receptor regulates ingestion, energy homeostasis, and body weight was what is the identity of the endogenous ligand for GUCY2C that represents a novel satiety factor? Valentino et al. found that the precursor to uroguanylin, prouroguanylin, is secreted after meals in both mice and humans (5). However, this left an important problem: prouroguanylin does not activate GUCY2C, it must be cleaved into uroguanylin, which then can activate that receptor. Where might this occur to result in satiation?

In a very clever experiment, Valentino et al. found that when either prouroguanylin or proguanylin were administered to a cell line expressing GUCY2C receptors, no increase in cGMP levels was observed (5). However, if prouroguanylin (but not proguanylin) was mixed with protein extracts from hypothalamus tissue, cGMP levels could be elevated. These experiments provided strong evidence to suggest that the hypothalamus is capable of converting prouroguanylin into active uroguanylin. Consistent with this conclusion, Valentino et al. found that GUCY2C receptors were expressed in the hypothalamus and that administration of a GUCY2C agonist directly into the CNS...
resulted in a reduction of food intake (5), providing tangible proof for a unique new component of the gut-brain signaling axis that controls metabolic homeostasis.

**Uroguanylin: a GI hormone different than others?**

Several GI hormones depend on unique posttranslational modifications as critical determinants of their actions. For example, glucagon-like peptide 1 (GLP-1) and glucose-dependent insulinotropic polypeptide (GIP) have a very short half-life, due to inactivation by the relatively ubiquitous protease dipeptidyl-peptidase 4 (DPP-4) (9). On the flip side of the DPP-4 equation is peptide tyrosine tyrosine (PYY), PYY is secreted as a 36–amino acid residue peptide and rapidly cleaved by DPP-4 into PYY3-36, which has been reported to control energy balance. A road to new therapies?

The discovery of uroguanylin as a satiety factor (5) adds yet another posttranslational strategy that the GI tract uses to sendafferent signals to the brain in order to maintain metabolic homeostasis. After meals, the hormonally inactive propeptide (prouroguanylin) appears to be secreted and transported via blood circulation to the CNS, in which it is converted to the active hormone, activates its cognate receptor, and suppresses food intake (Figure 1). This discovery impressively illuminates the many possibilities evolution has generated by which gut signals control systemic energy metabolism beyond classic endocrine mechanisms, such as modulating hormone secretion and receptor expression. It implies that, for reasons that are not obvious, it was advantageous for survival to find multiple alternative ways to fine tune the potency and the kinetics by which a complex array of endocrine signals could control energy balance.

**A road to new therapies?**

While the results of Valentino et al. (5) are extremely exciting and clinically significant, it is still too soon to declare victory in the fight to limit the obesity epidemic. There are several examples in which intriguing reports on novel peptide processing seemed to promise rapid breakthroughs...
comments

for the therapy of metabolic diseases but never delivered. In 2005, the “hormone” obestatin was reported as a previously overlooked by-product of preproghrelin cleavage with the ability to induce satiety in the CNS (12). Since then, scientists in the field have struggled to repeat the effects of obestatin on food intake or body weight seen in the previous report as well as to confirm its ability to bind and activate G protein–coupled receptor 39 (12).

With all of the GI hormones identified to date, it has turned out that their actions in the CNS are neither restricted to receptor activation in a single brain region nor limited to the modulation of a single physiological function. GLP-1 receptors, for example, are widely distributed throughout the brain and have been linked to visceral illness, anxiety, glucocorticoid secretion, and even promotion of neuronal survival (13, 14). As another example, ghrelin appears to act in key regions of the hippocampus, olfactory cortex, and ventral tegmental area to regulate memory, reward functions, and even mood (15–17), in addition to affecting memory, reward functions, and visceral illness. GLP-1 receptors mediate endocrine and anxiety responses to interoceptive and psychogenic stressors, J Neurosci 2003;23(15):6163–6170.


Behind an enteric neuron there may lie a glial cell

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The enteric nervous system (ENS) controls the gastrointestinal system. Enteric glia have long been regarded as the essential “glue” of the ENS. Now, however, two independent reports in this issue of the JCI provide compelling evidence that mouse enteric glia can also be neuronal precursors. These reports show that enteric glia give rise to neurons in vitro and that neurogenesis can be experimentally induced to occur in vivo in the adult mouse ENS. Unfortunately, glia do not constitutively replace neurons, and neurogenesis is not easily provoked. Although these new observations make it clear that clinical trials using glia to replace enteric neurons are more than premature, they are enticing for future research.

The enteric nervous system (ENS) is a neural crest–derived division of the autonomic nervous system that is essential for life (1).

It controls the gastrointestinal (GI) system and comprises a large number of neurons and glia that are organized into complex networks of interconnected ganglia distributed throughout the gut wall. Enteric neurons cluster into two plexi: the myenteric plexus, which is situated between the inner circular and outer longitudinal layers of the muscularis externa; and the submucosal plexus, which is located within the dense connective tissue between the muscularis externa and the mucosa.

One of the many functions of the GI system controlled by the ENS is motility, and thereby GI transit. As a result, aganglionosis of the bowel, whether congenital (as in Hirschsprung disease) or acquired (as in Chagas disease), leads to intestinal obstruction (2, 3). Nerve bundles, however, are present in the aganglionic segments of colon in individuals with Hirschsprung disease, meaning that nerves are not, by themselves, sufficient for GI transit (4). Nerve cell bodies and the complex microcircuits of the ENS, which uniquely enable it to control GI motility and secretion in the absence of CNS input, are required for normal GI tran-

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