Defective development and microcirculation of intestine in Npr2 mutant mice

Chizuru Sogawa-Fujiwara1, Atsuki Hanagata1, Yasuhiro Fujiwara2,3, Yukisato Ishida1, Hirotaka Tomiyasu4, Tetsuo Kunieda5, Hirofumi Nakatomi6 & Masatoshi Hori1

Intractable gastrointestinal (GI) diseases often develop during infancy. Our group previously reported that natriuretic peptide receptor B (NPR-B)-deficient Npr2<sup>slw/slw</sup> mice exhibit severe intestinal dysfunction, such as stenosis and distention, which resembles the dysfunction observed in Hirschsprung’s disease-allied disorders. However, the root cause of intestinal dysfunction and the detailed of pathophysiological condition in the intestine are not yet clear. Here, we report that the intestine of preweaning Npr2<sup>slw/slw</sup> mice showed bloodless blood vessels, and nodes were found in the lymphatic vessel. Additionally, the lacteals, smooth muscle, blood vessel, and nerves were barely observed in the villi of preweaning Npr2<sup>slw/slw</sup> mice. Moreover, intramuscular interstitial cells of Cajal (ICC-IM) were clearly reduced. In contrast, villi and ICC-IM were developed normally in surviving adult Npr2<sup>slw/slw</sup> mice. However, adult Npr2<sup>slw/slw</sup> mice exhibited partially hypoplastic blood vessels and an atrophied enteric nervous. Furthermore, adult Npr2<sup>slw/slw</sup> mice showed markedly reduced white adipose tissue. These findings suggest that the cause of GI dysfunction in preweaning Npr2<sup>slw/slw</sup> mice is attributed to defective intestinal development with microcirculation disorder. Thus, it is suggested that NPR-B signaling is involved in intestinal development and control of microcirculation and fat metabolism. This report provides new insights into intractable GI diseases, obesity, and NPR-B signaling.

The intestine has a layered structure and consisting of several tissues including the enteric nervous system (ENS) and interstitial cells of Cajal (ICC). The ENS controls intestinal motility, mucosal secretion, and sensory stimuli<sup>1</sup>. ICC are the pacemakers of the gastrointestinal (GI) tract and act as intermediaries between the ENS and smooth muscle<sup>2,3</sup>. Thus, disturbances of signals from the ENS and/or ICC affect motility and digestion/absorption<sup>4</sup>. Intractable GI diseases are rare, usually develop during infancy, and are often fatal. For example, in patients with Hirschsprung’s disease-allied disorders (HDAD)<sup>5–7</sup>, the ENS is present, yet distension and/or stenosis develop, causing reduced GI motility, a decline in GI transit, and excretory disorders<sup>8,9</sup>. Eventually, these defects lead to bacterial growth, resulting in enteritis and sepsis. However, the cause of GI intractable disease remains unclear.

NPR-B, also known as NPR2 or GC-B, is a transmembrane guanylyl cyclase receptor for the C-type natriuretic peptide (CNP), also known as NPPC. Activation of NPR-B by CNP binding synthesizes intracellular cyclic guanosine-3′,5′-monophosphate (cGMP)<sup>10–13</sup>. CNP also binds to natriuretic peptide receptor C (NPR-C, also known as NPR3 or GC-C)<sup>10,11</sup>, and CNP/NPR-B and/or NPR-C signals most likely act as local autocrine or paracrine factors in several tissues. Previous studies revealed that CNP/NPR-B and/or NPR-C signaling regulates contractility of the gastric antrum and large intestine<sup>14–17</sup>. It was also suggested that the CNP/NPR-B/cGMP signal may be related to ICC activity<sup>18–20</sup>, and vascular remodeling, and blood pressure<sup>21–28</sup>. Furthermore, CNP is also known to be related to adipocyte metabolism<sup>29,30</sup>.

<sup>1</sup>Veterinary Pharmacology, Graduate School of Agriculture and Life Sciences, The University of Tokyo, 1-1-1 Yayoi, Bunkyo-ku, Tokyo, Japan. <sup>2</sup>Institute for Quantitative Biosciences, The University of Tokyo, 1-1-1 Yayoi, Bunkyo-ku, Tokyo, Japan. <sup>3</sup>Graduate School of Human Health Sciences, Tokyo Metropolitan University, Minami-Osawa, Hachioji, Tokyo, Japan. <sup>4</sup>Veterinary Internal Medicine, Graduate School of Agricultural and Life Sciences, The University of Tokyo, 1-1-1 Yayoi, Bunkyo-ku, Tokyo, Japan. <sup>5</sup>Graduate School of Environmental and Life Sciences, Okayama University, Tsushima-naka, Kita-ku, Okayama, Japan. <sup>6</sup>Department of Neurosurgery, Graduate School of Medicine, The University of Tokyo, 7-3-1 Hongo, Bunkyo-ku, Tokyo, Japan. <sup>7</sup>email: chizuru.f.sogawa@gmail.com; y.fujiwara@iqb.u-tokyo.ac.jp
The short-limbed dwarfism (SLW) mouse has a mutation in Npr2, a gene encoding NPR-B, which results in the deletion of the intracellular domain of NPR-B. We previously reported that homozygous (Npr2<sub>slw/slw</sub>) mice frequently die pre-weaning due to GI dysfunction, such as abnormal retention of gastric milk, a stenosis-like phenotype in the terminal ileum and rectum, and distention of the GI tract by gas. In some cases, Npr2<sub>slw/slw</sub> mice showed no intestinal abnormality and were fully grown. It has also been recognized that the CNP/NPR-B signal increases cGMP and relaxes the stomach and large intestine. These observations indicate the important role of NPR-B in the physiological functions of the GI tract.

The CNP/NPR-B signal evidently regulates vital physiological functions in the GI tract. However, the way that NPR-B deficiency causes GI dysfunction in Npr2<sub>slw/slw</sub> mice has not been demonstrated conclusively. Thus, elucidation of the mechanism underlying GI dysfunction in Npr2<sub>slw/slw</sub> mice based on the detailed pathophysiological observations will provide valuable insights into GI diseases as well as for understanding CNP/NPR-B signaling.

### Results

#### Defective intestinal development in preweaning Npr2<sub>slw/slw</sub> mice.

In most preweaning Npr2<sub>slw/slw</sub> mice, abnormal distention was observed in the GI tract (Fig. 1a and Supplementary Fig. 1a). From the time of birth to approximately postnatal day 5 (P5), the small intestine (SI) of control mice, filled with yellow meconium, was covered with multi-branched blood vessels, whereas the amount of blood in blood vessels of Npr2<sub>slw/slw</sub> mice was evidently reduced (Fig. 1a, P0 and Supplementary Fig. 1b, P2 arrowheads). At approximately P7 to weaning age, the SI of control mice was filled with milk or fed bolus, and mesenteric blood vessels were more visible and covered with adipose tissue (Fig. 1a, control P8 and P18). However, in Npr2<sub>slw/slw</sub> mice, blood in mesenteric blood vessels and adipose tissue was barely observed (Fig. 1a, Npr2<sub>slw/slw</sub> P8 and P18 arrowheads). In some cases, SI was filled with viscous bubbles, discolored meconium, or only gas (Supplementary Fig. 1b, P10, P11, and P22). The nerves, blood vessels and lymphatic vessels in the mesentery were present in Npr2<sub>slw/slw</sub> mice, as seen in control mice (Fig. 1b i and and Supplementary Fig. 2a). Enteric neurons in the intestinal wall was also present in Npr2<sub>slw/slw</sub> mice, as seen in the control (Fig. 1b ii). In Npr2<sub>slw/slw</sub> mice, blood vessels in the intestinal wall were disrupted (Fig. 1b iii). Small vessels were observed in lymphatic vessels, and LYVE-1-positive macrophages were slightly larger in Npr2<sub>slw/slw</sub> mice than in control mice (Fig. 1b iv). Furthermore, a bifurcated region of lacteals was barely observed (Fig. 1b v). Although villi were formed, their internal contents were defective (Fig. 1b vi), and smooth muscle in villi was barely visible in preweaning age mice (Supplementary Fig. 2b). In contrast, control mice showed branched blood vessels but no lymphatic nodes, and development of those tissues was recognized, albeit at a younger age than Npr2<sub>slw/slw</sub> mice (Fig. 1b iii to vi, and Supplementary Fig. 2b). Moreover, a border of stenosis and distention, enteric neurons, was clearly recognized in the intestinal wall of Npr2<sub>slw/slw</sub> mice (Supplementary Fig. 2a, magnified image). Meconium and fed bolus were either liquid or absent in the large intestine (LI), and the LI was distended with gas in Npr2<sub>slw/slw</sub> mice (Fig. 1c and Supplementary Fig. 1c). In order to visualize adipocytes, we performed Oil Red-O staining of SI, and found intensely stained adipocytes along the mesenteric blood vessels in control mice (Fig. 1d). In contrast, in Npr2<sub>slw/slw</sub> mice SI, adipocytes were barely observed (Fig. 1d). Furthermore, in some cases, Npr2<sub>slw/slw</sub> mice showed cobweb-like microtubular structures filled with Oil Red-O stained microgranules (Supplementary Fig. 2c and d). Sexual differences were slightly skewed towards males (Fig. 1e); about 74% of Npr2<sub>slw/slw</sub> mice developed severe GI dysfunction before P20 (Fig. 1f).

We examined the health condition of Npr2<sub>slw/slw</sub> mice that exhibited severe GI disorder. Urinalysis showed a high index for ketone body and gravity (Fig. 2a, b and Supplementary Fig. 3a), and urine of Npr2<sub>slw/slw</sub> mice was yellowish (Supplementary Fig. 3b). This indicates that Npr2<sub>slw/slw</sub> mice did not absorb enough nutrients and were dehydrated. In some cases, Npr2<sub>slw/slw</sub> mice showed a higher protein index than control mice (Supplementary Fig. 3a), indicating renal dysfunction. The levels of triglycerides in the blood were not different between Npr2<sub>slw/slw</sub> and control mice (Fig. 2c). We also measured CNP levels in the blood, but no differences were observed between Npr2<sub>slw/slw</sub> and control mice (Fig. 2d). Electrolyte levels in the blood were not different between Npr2<sub>slw/slw</sub> and control mice at preweaning ages (Fig. 2e, f).

We assessed the morphology of regions of stenosis-like conditions such as the antrum, ileum terminal, and rectum. We compared the histological morphology of these regions between control and Npr2<sub>slw/slw</sub> mice at P10. From stomach to duodenum, gastric distention and atrophy were seen in some Npr2<sub>slw/slw</sub> mice (Fig. 3a stomach and Supplementary Fig. 4a stomach). In the regions before and behind the terminal ileum, the lumen of the...
Figure 2. Health condition in preweaning Npr2<sup>slw/slw</sup> mice. (a) Urinalysis of ketone body and (b) gravity. Control (n = 12) and Npr2<sup>slw/slw</sup> mice (n = 9). (c) Triglyceride level in the blood of control (n = 10) and Npr2<sup>slw/slw</sup> mice (n = 10). (d) CNP level in the blood of control (n = 19) and Npr2<sup>slw/slw</sup> mice (n = 21). (e) Electrolyte levels in the blood in P1 to P10 mice. Control (n = 8) and Npr2<sup>slw/slw</sup> mice (n = 7). (f) Electrolyte levels in the blood in P11 to P22. Control (n = 10) and Npr2<sup>slw/slw</sup> mice (n = 10). Bars indicate standard deviation (± SD).
terminus of control mice remained unconnected. In contrast, the distal part of ileum, cecum, and proximal colon of Npr2slw/slw mice were distended by gas, whereas the terminal ileum remained closed (Fig. 3a ileum terminal and Supplementary Fig. 4a ileum terminal). From the rectum to the anus, there were no morphological differences between control and Npr2slw/slw mice (Fig. 3a rectum and Supplementary Fig. 4a rectum). Additionally, Npr2slw/slw mice showed immature villi and mucosal epithelium in SI and LI (Fig. 3b), and intestinal edema in the distention region of the LI (Fig. 3b arrows head). We further examined the localization and morphology of enteric neurons by immunohistochemical staining of the antrum, ileum, and proximal colon using antibodies against PGP9.5, a pan-neuronal marker, Sox10, an enteric glial marker, and HuC/HuD, a ganglion cell marker. No differences were found in neuronal and enteric glial cells. However, ganglion cells appear to be slightly larger in Npr2slw/slw mice (Fig. 3c and Supplementary Fig. 4c). Additionally, western blot analysis confirmed that expression levels of PGP9.5, Sox10, and HuC/HuD were not significantly different between control and Npr2slw/slw mice, except for HuC/HuD in the colon, which were higher in Npr2slw/slw mice (Fig. 3d).

**Morphology of ICC and enteric neurons in preweaning Npr2slw/slw mice.** We examined the new-born status of ICC and enteric neurons, because in Npr2slw/slw mice, milk was seen in the stomach. To examine whether enteric neurons and ICC are affected in Npr2slw/slw mice, we comprehensively analyzed morphology of enteric neurons and ICC. In Npr2slw/slw mice at P0, around cardia, CD117-positive ICC was organized in a fan-like direction in both control and Npr2slw/slw mice (Fig. 4a). Although enteric neural crest-derived cells are known to migrate from the upper to lower regions of the GI tract, PGP9.5-positive enteric neurons and ICC were immature in antrum as compared with that of control mice, while they were normally present in the ileum to the rectum in both control and Npr2slw/slw mice (Fig. 4b and Supplementary Fig. 5a and b). Subsequently, at around 2 weeks of age, many Npr2slw/slw mice developed severe GI phenotype at this time. Enteric neurons and ICC were present in all regions in both mice (Fig. 5a) but enteric neurons were severely disrupted in the abnormally distended region in Npr2slw/slw mice (Fig. 5a, ileum). However, development and elongation of enteric neurons and ICC in the antrum and ileum terminal to rectum showed no differences between control and Npr2slw/slw mice (Fig. 5a). We measured the percent immunostained area for intramuscular ICC in the circular muscle layers (ICC-IM), terminal ileum, proximal colon, and rectum, and it was revealed that there were significantly fewer ICC-IM in Npr2slw/slw mice (Fig. 5b). In many cases, enteric neurons were present, not only ICC-IM but also ICC-MY, an Auerbach’s plexus layers, and ICC-IM in the longitudinal muscle layers of Npr2slw/slw mice were underdeveloped in LI compared with those of control mice (Supplementary Fig. 6a and b). We examined the reactivity of the intestinal smooth muscle using muscarinic and adrenergic agonists carbachol and noradrenaline. The result indicated that contractile and relaxant responses of smooth muscle to carbachol and noradrenaline were not different between control and Npr2slw/slw mice (Fig. 5d). Unexpectedly, western blot analysis of the whole layered GI tract showed that the expression level of CD117 in the LI was significantly higher in Npr2slw/slw mice than that in control mice, whereas no difference was shown in the stomach and ileum between control and Npr2slw/slw mice (Fig. 5e). Because CD117 is also a marker of mast cells, we examined mast cells of colon using FITC-avidin that is known to label mast cells. Whole-mount labeling revealed mesenteric mast cells were fewer in Npr2slw/slw mice than those in control. However, mucosal mast cells of Npr2slw/slw mice were often observed and apparently larger whereas those in control mice were rarely observed (Fig. 5f).

**Adult Npr2slw/slw mice.** Approximately 26% of Npr2slw/slw mice survived to adulthood, defined as mice over 6 months of age. Interestingly, gonadal white adipose tissue (GWAT) and mesenteric white adipose tissue of adult mice were evidently lower in Npr2slw/slw mice than those in the control mice (Fig. 6a and Supplementary Fig. 7a and b). The ratio of GWAT per body weight was significantly smaller in Npr2slw/slw mice than that in the control mice (Fig. 6b). Adipocytes were also smaller in Npr2slw/slw mice than those in the control mice (Fig. 6c), and the diameter of these adipocytes was significantly reduced in Npr2slw/slw mice compared with that in the control mice (Fig. 6d). The level of triglycerides in the blood was significantly lower in Npr2slw/slw mice than that in the control mice (Fig. 6e). Additionally, we frequently observed hypoplastic blood vessels with swollen areas in the SI of Npr2slw/slw mice ( Supplementary Fig. 7c). Significantly fewer branches of intestinal blood vessels were observed in Npr2slw/slw mice than in the control mice (Fig. 6f, g, and Supplementary Fig. 7d). Further, both normal and atrophied enteric neurons and ICC were observed in the SI of Npr2slw/slw mice (Fig. 6h and Supplementary Fig. 7e). The lymphatic vessel, bifurcation point of the lacteal, and lacteals with smooth muscle in villi were developed in both control and Npr2slw/slw mice (Fig. 6i). Additionally, they occasionally suffered obstructed defecation and debilitation (Fig. 6i and Supplementary Fig. 7f), and stools of aged Npr2slw/slw mice were smaller than those of the control mice (Fig. 6k). We thus analyzed the enteric neurons and ICC morphology of LI, and found that ICC including ICC-IM was developed in Npr2slw/slw and control mice (Fig. 6l for CML and LML, Fig. 6m for ICC-IM in the CML), while enteric neurons of Npr2slw/slw mice were atrophied in many cases (Fig. 6l, MY), but some regions showed normal enteric neurons (Supplementary Fig. 7g). Electrolyte levels in serum were not different between control and Npr2slw/slw mice (Fig. 6n).

**Discussion**

In our previous studies, we focused on smooth muscle function as a cause of GI dysfunction in Npr2slw/slw mice. However, the present study suggested that distention of Npr2slw/slw mice was not only attributed to stenosis but also to distension caused by defective development and circulation. It is likely that NPR-B-deficient Npr2slw/slw mice lost the ability to excrete waste, exchange gas, and absorb nutrients, which in turn caused viscus bubbles and corrosion of GI contents. However, although lacteals were not fully developed, the level of triglycerides in the blood was not significantly different compared with that in control mice at preweaning age. This may indicate that lipid absorption is not mediated by lacteal in lactation-period mice. Unlike lymphangiogenesis defective model
mice (including Prox1+/− mouse35 and vegfc deleted-mouse36), pups that showed intra-abdominal leakage of chylus, Npr2slw/slw mice did not exhibit such leakage. This is probably indicative of an intact lymphatic vessel itself.

CNP/NPR-B is known to regulate the vascular system. Given that the intestine of Npr2 slw/slw pups showed bloodless blood vessels, and that adults showed hypoplastic intestinal blood vessels, it strongly suggests that NPR-B plays an important role in the GI tract’s vascular system, including the bloodstream. Recent studies have suggested the important role of NPR-B and its ability to produce cGMP in the pericytes of microvessels reducing blood pressure27,28. Additionally, it was recently reported that CNP/NPR-C, rather than the CNP/NPR-B signal, plays an important role in angiogenesis and vascular remodeling23–26. This implies that NPR-B regulates blood pressure and blood flow, and in contrast, NPR-C regulates endothelial cells in the vascular system, which is involved in vascular remodeling.

Our comparative analysis between preweaning and adult mice clearly showed developmental delay of enteric neurons and/or ICC in the antrum and large intestine in Npr2slw/slw mice. This is probably the cause of abnormal retention in gastric milk and distention of the LI in pups. Additionally, high level of CD117 by western blot in Npr2slw/slw mice may be indicative of higher number of ICC-MY or enhanced proliferation of mucosal mast cells. However, Npr2slw/slw mice retained normal spontaneous motility and normal smooth muscle response by carbachol and noradrenaline ex vivo. Similarly, another lineage of NPR-B-mutant Npr2 cn/cn showed defective axonal bifurcation in spiral ganglion neurons and disorganized cochlear nuclei, and yet showed a normal sensitivity and response to sounds37.

This study also provides new findings regarding the maintenance of enteric neurons. Adult mice appeared healthy in many ways but tended to become debilitated earlier than controls. Indeed, atrophy of enteric neurons was observed in aged Npr2slw/slw mice. The hypoplastic blood vessels are probably related to neural maintenance,
Figure 5. Enteric neurons and ICC in 2 weeks mice and motility. (a) Whole mount immunostaining of tube specimens for PGP9.5 (green) and CD117 (magenta) were to identify enteric neurons and ICC, respectively in control (left) and Npr2slw/slw (right) mice. i: ileum; ce: cecum. Bars are 100 µm (images with short bars were captured using a 40 × objective lens; images with long bars were captured using a 20 × objective lens). Bars in terminal ileum are 1 mm. (b) Whole mount immunostaining of flat specimens for PGP9.5 (green) and CD117 (gray or magenta) were to identify enteric neurons and ICC in terminal ileum, proximal colon, and rectum. Upper row, Gray signal indicates ICC-IM in the circular muscle layer. Lower rows show the same position of upper row. enteric neurons (green) and ICC (magenta) were stacked from circular muscle layer to longitudinal muscle layer. Bars are 50 µm. (c) Columns indicate the area of ICC-IM in image field of terminal ileum, proximal colon, and rectum (n = 3 or 4 each for control and Npr2slw/slw mice, bars indicate ± SD). (d) Spontaneous rectal motility and response of smooth muscle by muscarinic and adrenergic agonists carbachol and noradrenaline. Triangles indicate the time of addition of 5 µM carbachol. Double triangles indicate the time of addition of 3 µM noradrenaline. Circles indicate the time of flushing with physiological salt solution (n = 3 each for control and Npr2slw/slw mice). CCh: carbachol; NA noradrenaline. (e) Comparison of protein level of CD117 by western blotting (n = 3 each for control and Npr2slw/slw mice). Columns indicate mean and bars indicate ± SD. a.u.: arbitrary unit. (f) Whole-mount labelling of colon flat specimens using FITC-avidin (green) was to identify mast cells. Serous side (left panels) and mucosal side (right panels) in control (n = 2) and Npr2slw/slw mice (n = 2). Mast cells are located within enclosed dashed circles or indicated by arrows. Bars are 100 µm.

and CNP/NPR-B may play an important role in the neurovascular unit in the GI tract. Furthermore, adult male Npr2slw/slw mice develop erectile dysfunction 8, probably also related to neurovascular function.

Overexpression of CNP specifically in endothelial cells or adipocytes (CNP-Tg mice) decreased mesenteric, epididymal, and subcutaneous fat, and prevented hypertrophy of adipocytes with a high-fat diet, whereas mice fed a standard diet showed no difference in adipocyte weight 25,30. Intriguingly, adult NPR-B-deficient Npr2slw/slw mice, in which lacteal developed normally, exhibited reduced perigonadal and mesenteric fat, both. This phenotype resembles that of CNP-Tg mice in which adipose tissues or adipocytes are reduced. This may imply an effect of the CNP/NPR-“C” function. Alternatively, the phenotype of Npr2slw/slw mice resembles that of CNP-Tg mice in which adipose tissues or adipocytes are reduced. This may imply an effect of the CNP/NPR-“C” function. Additionally, although GI dysfunction phenotypes have not been reported in other CNP/NPR-B mouse strains, it remains unclear why only Npr2slw/slw mice develop a unique GI phenotype that includes abnormalities of blood and lymphatic vessels.

Recent studies have suggested that CNP is related to inflammation. For example, CNP plasma concentrations increased in humans and rats with organ dysfunction caused by sepsis-related inflammation 38,39. However, CNP plasma levels were indistinguishable between control and Npr2slw/slw mice. The intestines of Npr2slw/slw mice did not turn red, and there were no signs of inflammation in Npr2slw/slw mice. Additionally, the level of electrolytes was indistinguishable between control and Npr2slw/slw mice. Indeed, the loss of NPR-B negatively regulates energy metabolism. Additionally, although GI dysfunction phenotypes have not been reported in other CNP/NPR-B mouse strains, it remains unclear why only Npr2slw/slw mice develop a unique GI phenotype that includes abnormalities of blood and lymphatic vessels.

In conclusion, our findings will help to understand the role of NPR-B in the GI tract and its microcirculatory system. Moreover, the observations of this study could be a clue to the cause of obesity and cryptogenic GI intractable disease.

Methods

Animal. An inbred SLW strain (Npr2slw: white coat) with a mixed genetic background of ddY (white coat) and C57BL/6 J (black coat) was obtained from the National Institute of Radiological Sciences (Chiba, Japan). To generate a congenic line C57BL/6J-Npr2slw (B6-Npr2slw), the inbred strain was backcrossed to the C57BL/6JSlsc (Japan SLC, Inc., Shizuoka, Japan) over seven generations, and subsequently heterozygous littermates were mated. An F1 SLW line was created from a mating between heterozygous inbred Npr2slw and B6-Npr2slw mice. An F2 SLW line was created from a mating between F1 heterozygotes of inbred and C57BL/6JSlsc mice. In this study, we used inbred, F1, and F2 lines, all of which showed the same phenotype. All mice were maintained under a standard 12-h light/dark condition. Either heterozygotes or wild-type mice were used for comparison (referred to as controls); homozygotes are referred to as Npr2slw. Followed by injecting an excess amount of a combined anesthetic containing 7.5% Domitor, 8% Midazolam and 10% Butorphanol tartrate in saline subcutaneously, all specimens were prepared.

All animal experiments were carried out in accordance with institutional guidelines regarding animal care and handling, and the experimental protocol was approved by the Institutional Animal Care and Use Committee of the University of Tokyo (Approval number P17-271H02).

Intestinal pharmacological response study. A pharmacological response study was performed as described previously 9, with some modifications. Briefly, the rectum of control and Npr2slw mice was collected and placed in physiological salt solution (PSS). Rectal strips with a whole layer were suspended longitudinally in 20 mL of PSS at 37 °C in an atmosphere containing 95% O2 and 5% CO2. After equilibration, each strip was exposed to 65.4 mmol potassium solution to obtain a stable contractile response. Subsequently, carbachol and noradrenaline were added to PSS to reach final concentrations of 5 µM and 3 µM, respectively. Contractile and relaxant movement profiles were recorded.

Quantification of protein expression level. Expression levels of marker proteins in the mouse GI tissue samples were measured by a Simple Western assay using a Jess (ProteinSimple, Santa Clara, CA, USA). Stomach
(corpus to antrum), ileum (distal region), and colon (proximal region) were homogenized with RIPA buffer (182–02,451, Fujifilm Wako Pure Chemical Corporation, Osaka, Japan) containing a protease inhibitor cocktail (P8340, Sigma Aldrich, MO, US), and loading samples were prepared using a 10× sample buffer (042–195, (182–02,451, Fujifilm Wako Pure Chemical Corporation, Osaka, Japan) for CD117/c-kit. Chemiluminescent signals were detected on a Jess Separation Capillary Cartridge (SM-W004, ProteinSimple). Primary antibodies used were rabbit polyclonal anti-PGP9.5 antibody (ADI-905–520-1, ENZO Life Sciences, Farmingdale, NY, US) for enteric nervous, intestinal glial cells, and goat polyclonal anti-CD117/c-kit antibody (AF1356, R&D systems) for ICC. Antibodies were diluted in an antibody diluent buffer (042–203, ProteinSimple) that was also used as a blocking buffer.

Whole mount immunofluorescence and fluorescence staining. For tube specimen, whole GI tissue was removed mesenterly and fixed in Bouin’s solution (85% saturated picric acid, 3.7% formaldehyde [10% 37% formaldehyde], and 5% glacial acetic acid) at room temperature (RT) for 2 h, and then immersed in 50% ethanol for 2 days. The specimens were trimmed, dehydrated and embedded in paraffin. Six-µm-thick sections were cut, placed on glass slides, and subjected to hematoxylin and eosin (HE) staining.

For immunostaining with diaminobenzene (DAB), antigen retrieval was performed by incubating in 10 mmol/L sodium citrate (pH 6.0) that contained 0.05% Tween-20 for 15 min at 95 °C, and gradually cooling to RT. The sections were washed in distilled water, immersed in TBS (50 mmol Trís–HCl, 150 mmol NaCl, and 2 mmol KCl), and then incubated in peroxidase-blocking solution (3% H2O2 in methanol) for 10 min at RT, followed by washing three times in TBS for 5 min at RT. The sections were then blocked with a non-specific blocking reagent (X0909, Dako, Glostrup, Denmark) for 10 min at RT, and incubated 2 h with the primary antibody at RT. The antibodies used were rabbit polyclonal anti-PGP9.5 antibody (ENZO Life Sciences, 1:200 dilution) for enteric nervous, mouse monoclonal anti-Huc/HuD antibody (A21271, Thermo Fisher, Waltham, MA, USA) for the ganglion cells, mouse monoclonal anti-Sox10 antibody (sc0365692, Santa Cruz Biotechnology, Santa Cruz, CA, USA) for intestinal glial cells, and goat polyclonal anti-CD117/c-kit antibody (AF1356, R&D systems) for ICC. Antibodies were diluted in an antibody diluent buffer (042–203, ProteinSimple) that was also used as a blocking buffer. Secondary antibodies used were anti-rabbit IgG antibody (042–206, ProteinSimple) for PGP9.5, anti-mouse IgG antibody (042–205, ProteinSimple) for HuC/HuD and Sox10, and anti-goat IgG antibody (705–035-147, Jackson ImmunoResearch Inc., West Grove, PA, USA) for CD117/c-kit. Chemiluminescent signals were detected using luminol-S (043–311, ProteinSimple) and peroxide (043–379, ProteinSimple). Data were analyzed using a software Compass for SW 4.0 (ProteinSimple).

Histology and immunostaining. Whole GI tissue was removed mesentry and fixed in Bouin's solution (85% saturated picric acid, 3.7% formaldehyde [10% 37% formaldehyde], and 5% glacial acetic acid) at room temperature (RT) for 2 h, and then immersed in 50% ethanol for 2 days. The specimens were trimmed, dehydrated and embedded in paraffin. Six-µm-thick sections were cut, placed on glass slides, and subjected to hematoxylin and eosin (HE) staining.

For immunostaining with diaminobenzene (DAB), antigen retrieval was performed by incubating in 10 mmol/L sodium citrate (pH 6.0) that contained 0.05% Tween-20 for 15 min at 95 °C, and gradually cooling to RT. The sections were washed in distilled water, immersed in TBS (50 mmol Tris–HCl, 150 mmol NaCl, and 2 mmol KCl), and then incubated in peroxidase-blocking solution (3% H2O2 in methanol) for 10 min at RT, followed by washing three times in TBS for 5 min at RT. The sections were then blocked with a non-specific blocking reagent (X0909, Dako, Glostrup, Denmark) for 10 min at RT, and incubated 2 h with the primary antibody at RT. The antibodies used were rabbit polyclonal anti-PGP9.5 antibody (ENZO Life Sciences, 1:200 dilution) for enteric nervous, mouse monoclonal anti-Huc/HuD antibody (Life Technologies Corporation, 1:100 dilution) for the ganglion cells, and mouse monoclonal anti-Sox10 antibody (Santa Cruz Biotechnology, 1:20 dilution) for intestinal glial cells. After incubation, the sections were washed with TBS three times for 5 min at RT and incubated with the secondary antibody for 1 h at RT. The secondary antibodies used were: ENVISION + rabbit HRP (K4002, Dako) for PGP9.5 and ENVISION + mouse HRP (K4000, Dako) for HuC/HuD and Sox10. After washing three times with TBS for 5 min at RT, the sections were stained with DAB (K3468, Dako) for 1–2 min at RT and then washed with distilled water. The nuclei were stained with hematoxylin.

Whole mount immunofluorescence and fluorescence staining. For tube specimen, whole GI tissue with mesentry was fixed in 10% buffer formalin overnight, and then immersed in TBS for overnight. The specimens were trimmed and washed with TBS. Subsequently, specimens were incubated in peroxidase-blocking solution for bleaching and demembranation for 30 min at RT. Followed by washing with TBS three times for 30 min at RT. The specimens were incubated with non-specific blocking reagent for 30 min at RT, and then reacted with the primary antibodies for 1–2 days at 4 °C on a shaker. The primary antibodies used were: rabbit polyclonal anti-PGP9.5 antibody (ENZO Life Sciences, 1:200 dilution) for enteric nervous, goat polyclonal anti-CD117/c-kit antibody (AF1356, R&D systems, 1:100 dilution) for ICC, rabbit polyclonal anti-alpha smooth muscle actin (aSMA) antibody (ab5694, Abcam, Cambridge, UK, 1:100 dilution) for smooth muscle, goat poly-
clonal LYVE-1 antibody (AF2125, R&D systems, 1:100 dilution) for lymphatic vessel, mouse monoclonal anti-PECAM antibody (sc-376764, Santa Cruz, CA, US, 1:20 dilution) for vascular endothelial cells, and FITC-avidin (434411, Thermo Fisher, 1:1,000 dilution) for mast cells. Each specimen was washed with TBS three times for 30 min at RT, and then incubated with the secondary antibodies overnight at 4 °C on a shaker in a dark condition. The secondary antibodies used were: donkey rabbit IgG H&L Alexa flour 488 (A21206, Thermo Fisher, 1:500 dilution) for PGP9.5 and aSMA, donkey goat IgG H&L 594 (A11001, Thermo Fisher, 1:500 dilution) for PECAM, and donkey goat IgG H&L Alexa flour 350 (A21081, Thermo Fisher, 1:500 dilution) for PECAM. Each specimen was washed with TBS three times for 30–60 min at RT, replaced with 50% glycerol in TBS on a glass bottom dish and covered with a cover glass.

For flat specimen, trimmed tube specimen as prepared above was cut longitudinally, immunostained, mounted with 50% glycerol TBS on a glass slide or glass bottom dish, and covered with a cover glass.

Oil Red-O staining. The whole GI tract with mesentery was fixed in 10% formalin overnight at 4 °C, and washed in tap water at RT for 1 h; then the specimens were trimmed and immersed in TBS until staining at 4 °C. The specimens were washed in doubly distilled water (DDW) for 3 min, immersed in 60% isopropyl alcohol for 2 min, and then immersed in filtered 60% Oil Red-O (4,049–1, Muto Pure Chemicals Co., Ltd, Tokyo, Japan) at 37 °C for 15 min. The specimens were washed in 60% isopropyl alcohol for 2 min and in DDW for 2 min.

Microscopy and acquisition. Stained images were acquired using BZ-X710 (Keyence, Osaka, Japan). HE and DAB stained sections were stacked and tiled. Wholemount fluorescent-immunostained and Oil Red-O stained whole-mount specimens were acquired using a BZ-H3XF/Sectioning Module. Unstained intestinal specimens were photographed using a stereo microscope (SZX16, Olympus).

Measurement of serum triglyceride. Triglyceride levels of serum were measured by DRI-CHEM SLIDE (470–03,701, Fujifilm Corporation, Tokyo, Japan) and DRI-CHEM auto tips (472–91,791, Fujifilm) using DRI-CHEM 7000 V (Fujifilm).

Measurement of serum electrolytes. Sodium, potassium, and chloride levels of serum were measured with an ion electrode method using a TBA-400FR Accute RX (Canon Medical Systems, Tochigi, Japan). The standard solution used was ISE CRS (J00700, Joko Co., Tokyo, Japan).

Measurement of plasma CNP concentration. Plasma CNP levels were measured with an NPPC ELISA kit for mouse (OKEH03207, Aviva Systems Biology, San Diego, CA, USA) according to the manufacturer’s instructions.

Urinalysis. Assay was performed using test papers (E-UR75, Eiken Chemical Co., Ltd, Tokyo, Japan). Urine was directly corrected by needle (29G SS-10M2913A, Terumo, Tokyo, Japan) under anesthesia. Each item was evaluated by dropping 10 µL of urine per item.

Statistical analysis. Data are expressed as a mean ± standard deviation (SD). A survival curve and dot plots were generated using GraphPad Prism7 software (GraphPad Software, San Diego, CA, USA). Area of ICC, enteric neurons, blood vessel and ICC-1M in image field and gonadal adipose white tissue (GWAT) diameter were calculated by Fiji software40. The statistical significance of differences in mean values was assessed by Student’s t-test.

Data availability
The data are available from the corresponding authors upon request.

Received: 17 April 2020; Accepted: 19 August 2020
Published online: 08 September 2020

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We thank Erina Inoue and Yuki Okada (Institute for Quantitative Biosciences, The University of Tokyo, Japan) and Satoshi Tsukamoto (National institute of Radiological Sciences, Chiba, Japan) for introducing the SWL mice to our mouse facility. We also thank Takako Yoshioka (Department of Pathology, National Center for Child Health and Development, Tokyo, Japan) for histological expertise, and Tatsuro Nakamura (Department of Animal Radiology, Graduate School of Agricultural and Life Sciences, The University of Tokyo) for the helpful discussion. Youichiro Wada and Yoshihiro Urade (Isotope Science Center, The University of Tokyo) for supporting the manuscript. This work is supported by a Grant-in-Aid of the Japan Society for the Promotion of Science (JSPS) Fellows (17J04685).

Acknowledgements
Author contributions
C.S.F. conceived the study, maintained the SLW mice, performed histological experiments and immunostaining analysis, analyzed the data and wrote the manuscript. A.H. performed the pharmacology and ELISA experiments. Y.F. did western blotting analysis, analyzed the data and wrote the manuscript. Y.I. measured electrolytes in the blood. H.T. measured triglyceride in the blood. T.K. provided the SLW mice. H.N. supported morphological analysis and provided reagents. M.H. supervised the study.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary information is available for this paper at https://doi.org/10.1038/s41598-020-71812-2.

Correspondence and requests for materials should be addressed to C.S.-F or Y.F.

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