Botulinum toxin A complex exploits intestinal M cells to enter the host and exert neurotoxicity

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To cause food-borne botulism, botulinum neurotoxin (BoNT) in the gastrointestinal lumen must traverse the intestinal epithelial barrier. However, the mechanism by which BoNT crosses the intestinal epithelial barrier remains unclear. BoNTs are produced along with one or more non-toxic components, with which they form progenitor toxin complexes (PTCs). Here we show that serotype A1 L-PTC, which has high oral toxicity and makes the predominant contribution to causing illness, breaches the intestinal epithelial barrier from microfold (M) cells via an interaction between haemagglutinin (HA), one of the non-toxic components, and glycoprotein 2 (GP2). HA strongly binds to GP2 expressed on M cells, which do not have thick mucus layers. Susceptibility to orally administered L-PTC is dramatically reduced in M-cell-depleted mice and GP2-deficient (Gp2⁻/⁻) mice. Our finding provides the basis for the development of novel antitoxin therapeutics and delivery systems for oral biologics.
Botulinum neurotoxin (BoNT), which is produced by *Clostridium botulinum* and related species, is a potent metalloprotease toxin consisting of a large protein (~150 kDa) that binds neuronal cells. On entering the cytoplasm of these cells, it cleaves SNAREs (soluble N-ethylmaleimide-sensitive fusion protein attachment protein receptors) required for vesicle fusion, thereby inhibiting neurotransmitter release and causing paralysis. BoNTs are produced along with one or more neurotoxin-associated proteins (NAPs) that non-covalently associate with BoNT to form progenitor toxin complexes (PTCs). The NAPs include non-toxic non-hemagglutinin (NTNHA) and hemagglutinin (HA). HA is composed of three different subcomponents: HA1, HA2 and HA3 (alternatively, HA-33, HA-17 and HA-70, respectively). HA1 and HA3 have carbohydrate-binding activities.

Intestinal absorption of BoNT is essential for the onset of food-borne botulism. However, the invasion site(s) and mechanism of BoNT in vivo are largely unknown. Here we analyze the site(s) responsible for intestinal translocation of the type A1 BoNT.

![Figure 1](image-url)
However, M-cell-dependent antigen uptake process which are present in the FAE. These cells effectively bind and cells present in the FAE. Therefore, we focused on the M cells, data imply that L-PTC binds to, and is internalized by, specific localization to any sites in the intestinal tissue (Fig. 1c). These loop assays in mouse. L-PTC was selectively localized at the FAE is transported to their basolateral sides via the interaction of HA in this step. L-PTC, which makes the predominant contribution of the toxin, we assessed localization of PTCs by intraperitoneally administered each toxin to mice (Fig. 1b). When intragastrically administered, the toxicity of L-PTC was highest, ~2 orders of magnitude greater than that of M-PTC and M-PTC was more toxic than BoNT alone. By contrast, there was no significant difference among the toxicities of the intraperitoneally administered toxins. These results indicate that L-PTC makes the predominant contribution to the onset of food-borne botulism.

To determine the site(s) responsible for intestinal translocation of the toxin, we assessed localization of PTCs by in situ intestinal loop assays in mouse. L-PTC was selectively localized at the FAE that covering PPs, whereas M-PTC exhibited no such clear localization to any sites in the intestinal tissue (Fig. 1c). These data imply that L-PTC binds to, and is internalized by, specific cells present in the FAE. Therefore, we focused on the M cells, which are present in the FAE. These cells effectively bind and deliver luminal macromolecules to the cells of underlying mucosal immune system for the induction of intestinal immune responses17. However, M-cell-dependent antigen uptake process can be exploited by some pathogens18. Indeed, L-PTC, its NAPs (a complex of NTNHA/HA) and HA bound to Ulex europaeus lectin I (UEA-1) + M cells, and were then transported to their basolateral sides (Fig. 1d and Fig. 5b). By contrast, M-PTC exhibited minimal interaction with M cells. Thus, HA is the critical factor in the interaction with M cells. After a 3-h incubation, L-PTC was located on the basolateral side of the FAE and in CD11c+ dendritic cells (CD11c+ DCs), which are located in the sub-epithelial dome (Supplementary Fig. 1a). Using L-PTC reconstituted with Alexa Fluor 488-labelled BoNT and Alexa Fluor 568-labelled NAPs, we also observed that CD11c+ DCs, localized beneath the M cells harbouring L-PTC and captured both BoNT and NAPs, which were only partially co-localized (Supplementary Fig. 1b,c, Supplementary Movie 1). Most of the NAPs were dissociated from L-PTCs and located predominantly in M cells, whereas most of the BoNTs were located in CD11c+ DCs. This observation was consistent with the previous proposal that NAPs, which are associated with BoNTs in the luminal environment of the intestine, dissociate after crossing the intestinal epithelium16. It remains unclear what percentage of BoNT present in the sub-epithelial dome is trapped by CD11c+ DCs. In any case, these observations indicate that all the constituents of L-PTC, including BoNT and NAPs, were transcytosed across the M cells to the basolateral surface, and captured by CD11c+ DCs. Furthermore, in vivo, L-PTCs were taken up by M cells in the small intestine 1 h after oral administration (Supplementary Fig. 2). Collectively, these results suggest that M cells play an important role in the transepithelial delivery of L-PTC.

Results
L-PTC is taken up by M cells via HA. To compare the toxicities of BoNT/A1 alone, M-PTC and L-PTC, we intragastrically or intraperitoneally administered each toxin to mice (Fig. 1b). When intragastrically administered, the toxicity of L-PTC was highest, ~2 orders of magnitude greater than that of M-PTC and M-PTC was more toxic than BoNT alone. By contrast, there was no significant difference among the toxicities of the intraperitoneally administered toxins. These results indicate that L-PTC makes the predominant contribution to the onset of food-borne botulism.

HA recognizes the sugar chains present on GP2. Next, we attempted to identify the critical subcomponent of HA required for the interaction with GP2 and localization of L-PTC to M cells. Individual HA subcomponents and the HA2/HA3 core complex formed by HA2 and HA3 (ref. 3) exhibited little or no binding to GP2 and no localization to M cells. By contrast, whole HA reconstituted with all three subcomponents strongly bound GP2 and localized to M cells in a manner similar to L-PTC (Fig. 5a–c). These results indicate that integration of all the HA subcomponents is required for the interaction with GP2, as well as for the localization of L-PTC to M cells. The fact that fully assembled HA is required for the interaction with GP2, even though the single subcomponents of HA1 and HA3 exhibit carbohydrate-binding activities4, suggests that these carbohydrate-binding activities are insufficient, either because single HA subcomponents lack sufficient avidity for robust binding or because the isolated subcomponents lack a three-dimensional
protein structure formed by the HA1/HA2/HA3 assembly involved in recognition of GP2. On the other hand, the presence of galactose inhibited binding between L-PTC and GP2 (Supplementary Fig. 5a), and L-PTC bound deglycosylated GP2 less strongly than intact GP2 (Supplementary Fig. 5b). To confirm the role of the carbohydrate-binding activities in GP2 recognition, we generated mutant HA complexes with an inactivated galactose-binding site (HA1:N285A/HA2/HA3, designated as N285A), an inactivated sialic acid binding site (HA1/HA2/HA3:R528A, designated as R528A) or both sites inactivated (HA1:N285A/HA2/HA3:R528A, designated as N285A/R528A) 7,25,26. The N285A complex exhibited dramatically reduced binding to GP2 and weak localization to M cells. The R528A complex also exhibited reduced binding to GP2 and weak localization to M cells, relative to the wild-type HA complex. The double-mutant complex, N285A/R528A, exhibited neither binding to GP2 nor localization to M cells (Fig. 5d,e). Furthermore, the HA complex bound to neither mGP2 (Fig. 5f) nor uromodulin (Supplementary Fig. 5c) expressed in a bacterial expression system yielding completely non-glycosylated proteins. These results indicate that the binding of HA complex to GP2 is mediated by the carbohydrate-binding activities of HA subcomponents, mainly the binding of hexavalent galactose-binding sites of HA1 to galactose moieties in sugar chains present on GP2.

**Discussion**

The observations reported here provide evidence that the BoNT/A1 complex exploits M cells to enter the host and exert neurotoxicity. Several studies have reported that experimentally purified BoNT/A1 and its heavy chain domain cross the intestinal epithelial barrier by transcytosis 27–31, and they preferentially recognize a subset of enteroendocrine cells32. However, BoNT is

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**Figure 2 | M cells are the major sites at which L-PTC breaches the intestinal epithelial barrier.** (a,b) Mice were treated intraperitoneally with anti-RANKL antibody or control rat IgG. (a) Alexa Fluor 568-labelled L-PTC (red) was injected into ligated mouse intestinal loops and incubated for 2 h, and FAE regions were stained with FITC-labelled UEA-1 (green) and iFluor 405-labelled phalloidin (blue). Scale bar, 20 μm. (b) L-PTC was intragastrically (1.0 pmol: 0.75 μg) or intraperitoneally (0.067 fmol: 50 pg) administered to mice treated with anti-RANKL antibody or rat IgG (intragastrically, n = 13 per group; intraperitoneally, n = 5 per group). Statistical analyses were performed with the log-rank test. NS, not significant. The data (a) are representative of two independent experiments.
Figure 3 | Glycoprotein 2 (GP2) serves as a major endocytotic receptor for L-PTC in M cells. (a) Recombinant Fc proteins were incubated with Strep-Tactin Superflow agarose pre-bound to fully assembled HA (HA1/HA2/HA3). HA-bound proteins were analyzed by immunoblotting using HRP-labelled anti-human IgG antibody. (b) Alexa Fluor 488-labelled L-PTC (green) were injected into ligated mouse intestinal loops and incubated for 1 h; GP2, uromodulin and PrPC were stained with specific antibodies and Cy3-labelled secondary antibodies (red). L-PTC predominantly co-localized with GP2 in M cells (arrows). For three-dimensional imaging of co-localization of L-PTC with GP2, see Supplementary Movie 2. Co-localization of L-PTC with GP2, uromodulin or PrPC in the randomly selected M cells was counted (n = 15), and the data are expressed as the number of merged L-PTC vesicles divided by the total number of L-PTC vesicles. Error bars indicate s.d. Statistical analyses were performed with Student’s t-test. Scale bar, 10 μm. Data are representative of two (b) or three (a) independent experiments.

Figure 4 | GP2 promotes the endocytotic uptake of L-PTC. MDCK I monolayers were transiently transfected with mGP2, and seeded on transwells. Monolayers were apically treated with L-PTC (200 nM) for 40 min at 4 °C (upper panels) or 1 h at 37 °C (lower panels). L-PTC was labelled with anti-L-PTC antibody and Alexa Fluor 488-labelled anti-rabbit IgG antibody (green), and GP2 was labelled with anti-GP2 antibody and Cy3-labelled anti-rat IgG antibody (red). Endocytotic vesicles of L-PTC in GP2-negative (GP2 −) or GP2-positive (GP2 +) cells were counted (GP2 − cells, 72 cells; GP2 + cells, 35 cells), and the data are expressed as total L-PTC vesicles per number of cells. Error bars indicate s.d. Statistical analyses were performed with Student’s t-test. L-PTC co-localized with GP2 in GP2 + cells (arrows). Scale bar, 10 μm. Data are representative of two independent experiments.
Figure 5 | Binding of HA with GP2 is mediated by the carbohydrate-binding activities of HA. (a,b) Alexa Fluor 568-labelled HA subcomponents or reconstituted HA (Alexa Fluor 568-labelled HA2/HA3 or HA1/Alexa Fluor 568-labelled HA2/HA3; red) were injected into ligated mouse intestinal loops and incubated for 2 h; FAE regions were stained with FITC-labelled UEA-1 (green). (c) Recombinant Fc proteins (mGP2 and hGP2) were incubated with Strep-Tactin Superflow agarose pre-bound to HA1, HA3, HA2/HA3 core complex or HA (HA1/HA2/HA3). HA-bound proteins were analyzed by immunoblotting using HRP-labelled anti-human IgG antibody. (d) Reconstituted WT HA (HA1/Alexa Fluor 568-labelled HA2/HA3), mutant HA complex harbouring mutant HA1 (N285A, HA1N285A/Alexa Fluor 568-labelled HA2/HA3), mutant HA complex harbouring mutant HA3 (R528A, HA1/Alexa Fluor 568-labelled HA2/HA3R528A) and mutant HA complex harbouring mutant HA1 and HA3 (N285A/R528A, HA1N285A/Alexa Fluor 568-labelled HA2/HA3R528A; red) were injected into ligated mouse intestinal loops and incubated for 2 h; FAE regions were stained with FITC-labelled UEA-1 (green). (e) Recombinant mGP2-Fc protein was incubated with Strep-Tactin Superflow agarose pre-bound to HA (WT, N285A, R528A or N285A/R528A). HA-bound proteins were analyzed by immunoblotting using HRP-labelled anti-human IgG antibody. (f) GST-mGP2 and GST were incubated with Strep-Tactin Superflow agarose pre-bound to HA (WT). HA-bound proteins were analyzed by immunoblotting using anti-GST antibody and HRP-labelled anti-rabbit IgG antibody. Scale bar, 10 μm (a,b,d). Data are representative of two (a,d) or three (b,c,e,f) independent experiments.
present as PTCs associated with NAPs in intestinal fluids. Therefore, the mechanisms underlying the absorption of PTCs are relevant to the pathogenesis of food-borne botulism. We cannot exclude the possibility that L-PTC is also taken up from absorptive epithelium and/or enteroendocrine cells. However, our observation that susceptibility to orally administered L-PTC was dramatically reduced in mice treated with anti-RANKL antibody (that is, M-cell-depleted mice) indicates that M cells are the major site for intestinal translocation of L-PTC.

In this study, we showed that the interaction of HA in the L-PTC with GP2 on the M-cell surface mediates the entry of L-PTC in mice. A similar mechanism may occur in humans, given that GP2 is specifically expressed by human M cells and L-PTC strongly binds human GP2 recombinant protein (Fig. 3a). These observations raise a question: what directs L-PTC to target GP2 on M cells? Indeed, HA binds to various glycoproteins expressed on the apical surface of enterocytes such as mucin 1 (MUC1) and dipeptidyl peptidase-4 (DPP4), as well as those expressed on the surface of lymphoid cells but not M cells; these include programmed cell death-1 (PD-1), T-cell immunoglobulin mucin protein 3 (TIM3), CD24 and CD40 although the binding to CD40 is weak (Supplementary Fig. 6). Moreover, L-PTC bound to the apical surface of MDCK I monolayers regardless of whether GP2 was present (Fig. 4). These facts suggest that there must be an additional factor involved in the M-cell tropism of HA. Enterocytes are protected from luminal pathogens by the thick layer of mucus and the filamentous brush border glycocalyx (FBBG) on the tip of microvilli, both of which are composed of highly glycosylated mucins. The FAE region has a relatively thin mucus layer compared with other regions of the intestinal mucosa, and this feature tends to promote local contact of antigens and pathogens with the FAE surface. Indeed, our data showed that L-PTC was trapped in the mucus layer and did not reach the enterocytes at the villous epithelium in situ intestinal loop assays (Supplementary Fig. 7). Furthermore, the FBBG acts as a size-selective barrier to limit the access of particles >30 nm in diameter. Enteroctyes in the FAE region have the FBBG, whereas M cells do not. Collectively, these observations suggest that the mucus layer and FBBG that cover the intestinal region in situ intestinal loop assays (Supplementary Fig. 7). Furthermore, the FBBG acts as a size-selective barrier to limit the access of particles >30 nm in diameter. Enteroctyes in the FAE region have the FBBG, whereas M cells do not. Collectively, these observations suggest that the mucus layer and FBBG that cover the intestinal region have the FBBG, whereas M cells do not. Collectively, these observations suggest that the mucus layer and FBBG that cover the intestinal region (Supplementary Fig. 8). These observations suggest that HA translocated via M cells serves to disrupt the adherens junctions.

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**Figure 6 | L-PTC is taken up by M cells via the GP2–HA interaction.** (a) Gp2+/+ or Gp2−/− mice were inoculated intragastrically (1.33 pmol: 1.0 μg) or intraperitoneally (0.067 fmol: 50 pg) with L-PTC (intragastrically, n = 15 per group; intraperitoneally, n = 5 per group). Statistical analyses were performed with the log-rank test. (b) Alexa Fluor 568-labelled L-PTC (red) were injected into ligated mouse intestinal loops and incubated for 1h. Images from FAE regions were analyzed quantitatively using the MetaMorph software. The data are expressed as percentages of the level in Gp2+/mice. Images from FAE regions were analyzed quantitatively using the MetaMorph software. The data are expressed as percentages of the level in Gp2+/+ mice. Error bars indicate s.d. For each group, six different PPs obtained from three different mice were examined (Gp2+/+, 127 cells; Gp2−/−, 161 cells). Statistical analyses were performed with Mann–Whitney U-test. Scale bar, 10 μm. NS, not significant.
between M cells and enterocytes, thereby compromising the paracellular barrier in vivo. Further study is needed to confirm whether this disruption of the paracellular barrier contributes to BoNT absorption in vivo.

Uptake of L-PTC from M cells was not abolished completely in Gp2–/– mice (Fig. 6b). In this regard, HA engaged in a weak interaction with uromodulin and PrPα (in vitro binding assay (Fig. 3a), which could explain the remaining L-PTC uptake by M cells. It should be noted, however, that GP2 was expressed at a much higher level than uromodulin and PrPα (Supplementary Fig. 4). In addition, among these three ligands, L-PTC in the endocytic vesicles co-localized most frequently with GP2. Consistent with these notions, susceptibility to orally administered L-PTC was markedly reduced in Gp2–/– mice, suggesting that GP2-mediated endocytosis is the critical pathway for the translocation of L-PTC in the context of food-borne botulism. Further investigation will be needed to elucidate how BoNT deals with the immune surveillance in PPs and disseminates into the systemic circulation. Some BoNT serotypes (for example, serotypes E and F) cause food-borne intoxication even though they lack genes encoding HA proteins and produce only M-PTCs. Further studies will be required to elucidate the intestinal absorption mechanism of these M-PTCs.

Taken together, our findings shed light on the process of intestinal BoNT infection, heretofore largely unknown, which is essential. This knowledge could facilitate development of more effective therapies for this disease. Furthermore, techniques for M-cell-targeted delivery of vaccine antigens could greatly improve the efficacy of mucosal vaccines. Thus, exploiting the function of HA represents an attractive approach to development of novel delivery systems for vaccines and drugs.

Methods

Animals. BALB/c mice were purchased from SLC. GP2-deficient mice (Gp2–/– mice) were generated as described previously37,38, and then backcrossed into a BALB/c genetic background. By tissue staining with an anti-GP2 monoclonal antibody, we verified previously that the GP2 knockout abolishes GP2 expression on M cells.39 These mice were maintained under specific pathogen-free conditions. Female mice were used at the age of 8–12 weeks (Figs 1,3 and 5 and Supplementary Figs 1,2,4,7 and 8), 8 weeks (Fig. 2 and Supplementary Fig. 3) or 11–19 weeks (Fig. 6). Mice of similar ages were used in comparative analyses. The experimental protocols were approved by the Ethics Review Committee for Animal Experimentation at Osaka University.

Toxins and non-toxic components. Toxins and non-toxic components were purified from the culture fluid of C. botulinum type A strain 62A (ref. 8). Briefly, the organisms were cultured using a cellobiose-tube procedure, and the culture supernatant was collected and concentrated by 60% ammonium sulfate precipitation. After dialysis, toxin-containing solution was applied to a glycine-sulfate precipitation. After dialysis, toxin-containing solution was applied to a cellulose membrane (G2). The specimens were then washed with PBS, and subjected to confocal microscopy using a CSU21 or CSUX1 confocal scanner unit (Yokogawa) and an IX71 microscope (Olympus). The data were analyzed using the MetaMorph software (Molecular Devices).

Competition of toxicity. Mice were not fed for 4 h prior to administration, but had free access to water. Mice were inoculated intragastrically by gavage with 0.3 ml of 10 mM sodium phosphate buffer (pH 6.0) ± 0.1% gelatin containing various concentrations of toxins. Alternatively, mice were injected intraperitoneally with 0.3 ml of 10 mM sodium phosphate buffer (pH 6.0) ± 0.1% gelatin containing various concentrations of toxins. We used a toxin dose at which most mice demonstrated flaccid quadriplegia ~24 h after the administration of PTCs. Mice were observed for morbidity, mortality and survival time over the course of 3 weeks. The investigators were ‘blinded’ to group allocation both during the experiment and when assessing the outcome (Figs 2b,6a and Supplementary Fig. 3).

Preparation of HA proteins. Purified DNA from C. botulinum type A strain 62A was used as a template for the amplification of DNA encoding HA1, 2 and 3 by PCR. The amplified DNAs were inserted into pET-52b(+) (Invitrogen) and visualized by confocal microscopy using a CSU21 or CSUX1 confocal scanner unit (Yokogawa) and an IX71 microscope (Olympus). The data were analyzed using the MetaMorph software (Molecular Devices). For quantitative analysis of localization of L-PTC to M cells, Alexa Fluor 546-labelled L-PTC (20 pmol per 100 μl) was injected into ligated intestinal loops containing PPs and incubated for 1 h. Whole-mount specimens were stained with FITC-labelled UEA-1. Images were taken from FAE and analyzed using the MetaMorph software. The localization value was calculated as follows: (L-PTC area × average intensity of L-PTC in this area)/UEA-1 area.

Immunofluorescence staining. After ligated intestinal loop assays, whole-mount specimens were fixed with 4% paraformaldehyde for 30 min at room temperature, washed with PBS (pH8.0 and 0.75% NaCl) and then permeabilized for 5 min in 50 mM NH4Cl for 30 min at room temperature. After permeabilization with 0.5% Triton X-100, specimens were incubated with blocking buffer (2% BSA in PBS) for 1 h, followed by incubation with primary antibodies (anti-GP2 (ref. 21) (2.5 μg ml−1, Jackson Immunoresearch) or Alexa Fluor 488-labelled secondary antibody (10 μg ml−1, Invitrogen) for 2 h at room temperature. The specific staining of primary antibodies (anti-GP2, anti-uromodulin and anti-PrPα) was confirmed by comparison with an isotype control primary antibodies. In some experiments, specimens were stained for 2 h at room temperature with FITC or rhodamine-labelled UEA-1 (4.0 μg ml−1, Vector Laboratories), Alexa Fluor 568-labelled phallodin (Invitrogen), iFluor 405-labelled phallodin (Abcam) or Pacific green-labelled anti-CD11c antibody (Life Technologies). After immunofluorescence staining, specimens were mounted in ProLong Antifade (Invitrogen) and visualized by confocal microscopy using a CSU21 or CSUX1 confocal scanner unit (Yokogawa) and an IX71 microscope (Olympus). The data were analyzed using the MetaMorph software (Molecular Devices).

For mucin staining, whole-mount specimens were fixed with Carnoy’s solution (60% ethanol, 30% chloriform and 10% glacial acetic acid) for 1 h at room temperature, and then washed with a series of 100% ethanol, 70% ethanol, 40% ethanol and PBS. The specimens were incubated with blocking buffer for 1 h, and then incubated with anti-MUC2 antibody (2.0 μg ml−1, Abcam) for 2 h at room temperature. The specimens were washed with PBS, and then probed with Cy3-labelled secondary antibody and iFluor 405-labelled phallodin for 2 h at room temperature.

For quantitative analysis of localization of L-PTC to M cells, Alexa Fluor 568-labelled L-PTC (20 pmol per 100 μl) was injected into ligated intestinal loops containing PPs and incubated for 1 h. Whole-mount specimens were stained with FITC-labelled UEA-1. Images were taken from FAE and analyzed using the MetaMorph software. The localization value was calculated as follows: (L-PTC area × average intensity of L-PTC in this area)/UEA-1 area.

Treatment with anti-RANKL antibodies. The anti-mouse RANKL monoclonal antibody (clone IK22-5) was generated by immunizing Sprague Dawley rats with recombinant mouse RANKL and fusing immune lymph node cells with myeloma cells. This antibody was used for in vivo RANKL neutralization to transiently deplete M cells.40 Mice were injected intraperitoneally with 250 μg of anti-RANKL antibody every 2 days for 8 weeks. As a negative control, another group of mice was treated with a purified functional-grade isotype control rat immunoglobulin-G2a (IgG2a) antibody (ebioscience).

Comparison of toxicity. Mice were not fed for 4 h prior to administration, but had free access to water. Mice were inoculated intragastrically by gavage with 0.3 ml of 10 mM sodium phosphate buffer (pH 6.0) ± 0.1% gelatin containing various concentrations of toxins. Alternatively, mice were injected intraperitoneally with 0.3 ml of 10 mM sodium phosphate buffer (pH 6.0) ± 0.1% gelatin containing various concentrations of toxins. We used a toxin dose at which most control mice presented with flaccid quadriplegia ~24 h after the administration of PTCs. Mice were observed for morbidity, mortality and survival time over the course of 3 weeks. The investigators were ‘blinded’ to group allocation both during the experiment and when assessing the outcome (Figs 2b,6a and Supplementary Fig. 3).

Preparation of HA proteins. Purified DNA from C. botulinum type A strain 62A was used as a template for the amplification of DNA encoding HA1, 2 and 3 by PCR. The amplified DNAs were inserted into pET-52b(+) (Invitrogen) and visualized by confocal microscopy using a CSU21 or CSUX1 confocal scanner unit (Yokogawa) and an IX71 microscope (Olympus). The data were analyzed using the MetaMorph software (Molecular Devices).
For immunofluorescence staining, HA proteins were labelled with Alexa Fluor dye. The degrees of labelling: Strep-HA1, 1.4 mol dye per mol protein; FLAG-HA2, 0.9 mol dye per mol protein; Strep-HA3, 2.1 mol dye per mol protein.

**Preparation of Fc-fusion proteins.** To obtain expression vector for fusion proteins of mouse GP2, human GP2, mouse uromodulin (Umod) or mouse PrPC with the Fc segment of human IgG1 (mGP2-Fc, gp2-Fc, Umod-Fc and PrPC-Fc), complementary DNAs (cDNAs) corresponding to the extracellular domains of each protein were amplified by PCR using primer sets listed in Supplementary Table 1. The cDNA fragments were inserted into the pcDNA3 expression vector (Invitrogen), and then cultured for several days. The Fc-fusion proteins secreted were detected by immunoblotting using an anti-GST antibody purified with a GSTrap column (GE Healthcare) and used for pull-down assays.

**Enzyme-linked immunosorbent assay (ELISA).** Primer sets are listed in Supplementary Table 1. Samples were amplified with the Eco Real Time PCR System (Illumina), SYBR Green I and the tracer was injected into ligated mouse intestinal epithelium to measure. In the competition assay, L-PTC was pre-incubated with sugar chain for 2 h at 4 °C. Wells were washed three times with PBS (pH 7.4) overnight at 4 °C. Wells were washed three times with PBS (pH 6.0), blocked with 1% BSA in PBS and incubated with L-PTC for 1 h at room temperature. After washing, HRP-labelled anti-rabbit IgG antibody (diluted 1:10,000) and HRP-labelled anti-rabbit IgG antibody (diluted 1:1,000) for 2 h at room temperature. After washing, rabbit anti-BoNT polyclonal antibody (diluted 1:3,000) was added and plates were incubated for 1 h at room temperature.

**In situ paracellular flux assay.** Mice were not fed for 4 h prior to surgery, but had free access to water. During the procedure, mice were maintained under pentobarbital sodium anaesthesia. NAPs (NTNHA/HA, 500 pmol) and paracellular tracer (fixable FITC-dextran 10 K, D-10, 2.0 mg ml⁻¹, Invitrogen, 100 µl) or vehicle control (PBS) and the tracer were injected into ligated mouse intestinal loops and incubated for 2 h, and FAE regions were stained with rhodamine-labeled UEA-1. After immunofluorescence staining, the morphology of M cells and localization of paracellular tracer were observed by confocal microscopy.

**Quantitative reverse-transcription PCR (qRT-PCR).** Total RNA was isolated from FAE of BALB/c mouse using the RNeasy kit (Life Technologies) and subjected to reverse transcription using the PrimeScript kit (TAKARA BIO). The cDNA samples were amplified with the Eco Real Time PCR System (Illumina), SYBR premix ex Taq (TAKARA BIO) and the primer sets specific for mouse genes. Primer sets are listed in Supplementary Table 1.

**Enzyme-linked immunosorbent assay (ELISA).** Binding of L-PTC to recombinant mouse GP2-Fc was analyzed by enzyme-linked immunosorbent assay (ELISA). Plates were coated with recombinant mouse GP2-Fc protein at a final concentration of 2.5 µg ml⁻¹ in PBS (pH 7.4) overnight at 4 °C. Wells were washed three times with PBS (pH 6.0), blocked with 1% BSA in PBS and incubated with L-PTC for 1 h at room temperature. After washing, rabbit anti-BoNT polyclonal antibody (diluted 1:3,000) was added and plates were incubated for 1 h at room temperature. After washing, HRP-labelled anti-rabbit IgG antibody (diluted 1:10,000) was added, and the sample was incubated for an additional 1 h at room temperature. Plates were then treated with ABTS (Roche) and absorbance values (optical density: 405 nm) were measured. The competition assay, L-PTC was pre-incubated with sugar chain for 1 h at room temperature prior to ELISA.

**Glycopeptidase F treatment.** Recombinant mouse GP2-Fc was treated with Glycopeptidase F (PNGase F, TAKARA BIO) to remove N-linked glycan from GP2. Briefly, 3 µg of mGP2-Fc was incubated with Glycopeptidase F (5.0 µU) for 24 h at 37 °C in 1% (w/v) HEPES (Dojindo) at pH 7.4 (ref. 14). Cells were transiently transfected with vector encoding mouse GP2 using ScreenFect A (Wako) and seeded on Transwell pore filter (Costar). After 3 days, cells were apically treated with L-PTC. After various times, cells were fixed with 4% paraformaldehyde for 20 min and permeabilized with 0.5% Triton X-100 for 5 min, and then incubated with blocking buffer (5% BSA in PBS) for 1 h. The cells were incubated with anti-L-PTC (diluted 1:400) and anti-GP2 antibodies (1.0 µg ml⁻¹) and then probed with Alexa Fluor 488 (5.0 µg ml⁻¹) or Cy3 labelled secondary antibodies (1.5 µg ml⁻¹).

**Statistical analyses.** Student’s t-test was used to compare the binding of L-PTC with Fc-fusion proteins; the co-localization of L-PTC with GP2, uromodulin or PrPC in M cells and the endocytotic vesicles of L-PTC in MDCK I cells. The Mann–Whitney U-test was used to compare the localization of L-PTC to M cells in FAE. The Kaplan–Meier estimator method was used to draw survival curves, and the log-rank test was used to compare the survival rates. P values <0.05 were considered to be statistically significant.

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Author contributions

T.M., H.O., and Y.F. designed the research and analyzed the data. T.M. performed the majority of the experiments and the analyses. Y.S., M.Y., and S.A. provided recombinant materials and analyzed the data. H.Y. provided the anti-RANKL antibody. T.K. provided toxins and anti-BoNT antibodies. S.F. provided GP2-deficient mice. T.N. performed the qPCR assays and analyzed the data. K.H., S.F. and H.O. provided Fc-fusion proteins. T.M., H.O., K.H. and Y.F. co-wrote the manuscript. All authors participated in the discussion of the data and in production of the final version of the manuscript.

Additional information

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