Paneth Cell Secretion in vivo Requires Expression of Tmem16a and Tmem16f

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BACKGROUND AND AIMS: Paneth cells play a central role in intestinal innate immune response. These cells are localized at the base of small intestinal crypts of Lieberkuhn. The calcium-activated chloride channel TMEM16A and the phospholipid scramblase TMEM16F control intracellular Ca2+ signaling and exocytosis. We analyzed the role of TMEM16A and TMEM16F for Paneth cells secretion. METHODS: Mice with intestinal epithelial knockout of Tmem16a (Tmem16a<sup>−/−</sup>) and Tmem16f (Tmem16f<sup>−/−</sup>) were generated. Tissue structures and Paneth cells were analyzed, and Paneth cell exocytosis was examined in small intestinal organoids in vitro. Intracellular Ca2+ signals were measured and were compared between wild-type and Tmem16 knockout mice. Bacterial colonization and intestinal apoptosis were analyzed. RESULTS: Paneth cells in the crypts of Lieberkuhn from Tmem16a<sup>−/−</sup> and Tmem16f<sup>−/−</sup> mice demonstrated accumulation of lysozyme. Tmem16a and Tmem16f were localized in wild-type Paneth cells but were absent in cells from knockout animals. Paneth cell number and size were enhanced in the crypt base and mucus accumulated in intestinal goblet cells of knockout animals. Granule fusion and exocytosis on cholinergic and purinergic stimulation were examined online. Both were strongly compromised in the absence of Tmem16a or Tmem16f and were also blocked by inhibition of Tmem16a/f. Purinergic Ca2+ signaling was largely inhibited in Tmem16a knockout mice. Jejunal bacterial content was enhanced in knockout mice, whereas cellular apoptosis was inhibited. CONCLUSION: The present data demonstrate the role of Tmem16 for exocytosis in Paneth cells. Inhibition or activation of Tmem16a/f is likely to affect microbial content and immune functions present in the small intestine.

Keywords: Paneth Cells; Exocytosis; TMEM16A; Anoctamin 1; ANO1; TMEM16F; Anoctamin 6; ANO6; Goblet Cells; Secretion; Ca2+-Activated Cl− Channel; Phospholipid Scrambling

Introduction

Paneth cells have a central function in intestinal innate immune response.1–3 Paneth cells are located in the base of small intestinal crypts of Lieberkuhn and have defensive functions that include protection of stem cells in response to invading microbes and eradication of ingested pathogens. Together with other secretory cells, they clear pathogens from intestinal crypts.4 By means of secreted factors, they also regulate the composition and number of commensal intestinal bacteria.5 Paneth cells are filled with rather large granules that contain antimicrobial proteins/peptides, such as lysozyme, secretory phospholipase-A2, and defensins, and they also secrete cytokines to recruit immune cells.6 During Paneth cell metaplasia in inflammatory bowel disease or as a response to mucosal damage, the Paneth cell zone expands due to increase in cell size and cell number.7 It was shown that cholinergic stimulation confers enhanced protection in animals orally infected with virulent Salmonella enterica.8 Acetylcholine binds to basolateral M3 muscarinic receptors allowing adaptive immunity to helminth and bacterial infection.9 However, the mechanisms of luminal stimulation of Paneth cell secretion via life bacteria or lipopolysaccharide (LPS) are unclear.4 Paneth cells do not express Toll-like receptor 4 (TLR4), the receptor for LPS.10–12 However, LPS can also trigger immune response independent of TLR4 by binding to LPS-binding protein, which is then internalized by the host cell.13,14 It was shown that uptake of LPB/LPS triggers subsequent adenosine triphosphate (ATP) release.15

We recently reported for the first-time activation of intestinal mucus secretion by luminal and basolateral ATP.16 ATP activates mucus secretion via activation of purinergic P2Y2 receptors and release of Ca2+ to the apical submembraneous compartment. For efficient release of Ca2+ via inositol trisphosphate (IP3)–activated Ca2+ release channels (IP3R) located in the endoplasmic reticulum (ER), IP3R is tethered to the apical compartment by binding to the Ca2+-activated Cl− channel TMEM16A (anoctamin 1, ANO1).16,17 High local Ca2+ levels in the apical intracellular compartment support the exocytotic machinery in intestinal and airway goblet cells.18,19 Importantly, ATP-induced

Abbreviations used in this paper: ATP, adenosine triphosphate; CCH, carbachol; CF, cystic fibrosis; ER, endoplasmic reticulum; LPS, lipopolysaccharide; TLR4, Toll-like receptor 4; TUNEL, Terminal deoxynucleotidyl transferase dUTP nick end labeling; WT, wild type.
mucus secretion by secretory cells was compromised in both intestine and airways of Tmend16a knockout mice (Tmend16a−/−).18 As a consequence, mucus accumulated in intestinal and airway goblet cells of Tmend16a−/− mice. A similar but less pronounced mucus accumulation was also observed in Tmend16f−/− mice.20 Here, we demonstrate the role of Tmend16a and Tmend16f for exocytosis in Paneth cells and describe the underlying molecular mechanisms. The present findings are of clinical relevance, because many small molecules and natural or herbal compounds exist that either activate or inhibit both Tmend16a and Tmend16f.21,22

Methods

Animals

Generation of mice with intestinal epithelial-specific knockout of Tmend16a (Tmend16a−/−) or Tmend16f (Tmend16f−/−) and genotyping has been reported earlier.20,23

Intestinal Crypts Isolation, Organoids Culture, and Life Microscopy Imaging (Differential Interference Contrast)

Isolation of intestinal crypts and culturing of organoids were done as described by Mahe et al.24 and Altay et al.25 Organoids in Matrigel (Corning, Wiesbaden, Germany) were cultured on glass coverslips in media containing (GIBCO; Thermo Fisher, Scientific, Waltham, MA, USA) Advanced DMEM/F-12, L-glutamine, (4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid), penicillin/streptomycin, N2 supplement, B27 supplement, mouse recombinant EGF (Sigma-Aldrich, Merck KGaA, Darmstadt, Germany), Mouse Recombinant Noggin (PeproTech, Hamburg, Germany), Human Recombinant R-Spondin-1 (PeproTech), and N-acetyl-L-cysteine (Sigma-Aldrich, Merck KGaA). Three- to six-day-old organoids were mounted into a chamber overlaid with 100 μL Ringer solution (NaCl 145 mmol/L; KH2PO4 0.4 mmol/L; K2HPO4 1.6 mmol/L; glucose 5 mmol/L; MgCl2 1 mmol/L; Ca2+1.33 mmol/L). Paneth cells were observed using an Axiovert Observer microscope (Carl Zeiss Microscopy Deutschland GmbH, Oberkochen, Germany) using differential interference contrast. Pictures are taken every 10 seconds using the ZEN software (Carl Zeiss Microscopy Deutschland GmbH). Secretion was stimulated by ATP (100 μM; Sigma-Aldrich, Merck KGaA) and subsequently with carbachol (CCH, 10 μM; Sigma-Aldrich, Merck KGaA). Granule secretion was analyzed according to Yokoi et al.1 using the ZEN software.

Histology and Immunohistochemistry of Tmend16a, Tmend16f, and Lysozyme

Mouse duodenum, jejenum, and ileum were fixed with 4% paraformaldehyde, 0.2% picric acid, and 3.4% sucrose overnight, dehydrated, and embedded in paraffin. The paraffin-embedded tissues were cut at 4 μm on a rotary microtome (Leica Mikrotom RM 2165, Wetzlar, Germany). The sections were dewaxed and rehydrated. For histology or mucus analysis, sections were stained according to standard hematoxylin/eosin or periodic acid–Schiff protocols and examined by light microscopy. For immunohistochemistry, sections were cooked in Tris/ethylene diaminetetraacetic acid (pH8.5, TMEM16A or TMEM16F staining) or citrate buffer (pH 6, for lysoyme staining) for 15 minutes and permeabilized and blocked with 0.04% Triton X-100 and 5% bovine serum albumin for 30 minutes at 37 °C. Sections were incubated with primary antibodies against mouse TMEM16A antigen MEECAPGQCLMLCQIQ1 (Davids Biotechnologie GmbH, Regensburg, Germany), mouse TMEM16F antigen KREKYLYQLSHHELKDLT (Davids Biotechnologie GmbH) or lysoyme (PA5_16668; Invitrogen, Thermo Fisher, Scientific) in 0.5% bovine serum albumin and 0.04% Triton X-100 overnight at 4 °C and subsequent with a secondary goat anti-rabbit Alexa 488 or goat anti-rabbit Alexa 546 IgG (Invitrogen, Thermo Fisher, Scientific) for 1 hour at 37 °C. Sections were counterstained with Hoe33342 (Sigma-Aldrich, Merck KGaA). Immunofluorescence was detected using an Axiovert Observer microscope equipped with ApoTome2 and ZEN software.

Apopotic Cell Death in Mouse Intestine

Terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) was performed in 4% paraformaldehyde, 0.2% picric acid, and 3.4% sucrose fixed and in paraffin-embedded murine jejunum and ileum. For TUNEL assay, the DeadEnd Colorimetric TUNEL System (Promega, Mannheim, Germany) was used according to manufacturer’s instructions and analyzed with ImageJ (NIH, USA).26

Detection of Bacteria in Mouse Jejunum and Ileum

Gram-positive and Gram-negative bacteria were detected using Gram Stain Kit (Abcam, Amsterdam, the Netherlands). Briefly, after deparaffinization and hydration, paraffin-embedded jejunum or ileum was incubated in Gentian violet solution and rinsed in water. Slides were incubated in Lugol’s iodine solution and rinsed with water. Slides were decolorized with Gram decolorizer and then incubated in carbol fuchsin. After washing, the tissue elements were counterstained with tartrazine solution. The number of bacteria in cross-sections was analyzed using ImageJ.26

Measurement of Intracellular Ca2+ in Paneth Cells

Intestinal organoids on glass coverslips were loaded with 2 μM Fura-2/AM and 0.02% Pluronic F-127 (Invitrogen, Thermo Fisher, Scientific) in ringer solution (NaCl 145 mmol/L; KH2PO4 0.4 mmol/L; K2HPO4 1.6 mmol/L; glucose 5 mmol/L; MgCl2 1 mmol/L; Ca2+1.33 mmol/L) for 1 hour at room temperature. The agonists ATP and CCH were applied subsequently to increase intracellular Ca2+ concentrations. Ca2+ increase by application of CCH and ATP in reverse sequence was not significantly different. Fluorescence was detected at 37 °C using an inverted microscope (Axiovert S100, Carl Zeiss Microscopy Deutschland GmbH) and a high-speed polychromator system (VisiChrome, Puchheim, Germany). Fura-2 was excited at 340/380 nm, and emission was recorded between 470 nm and 550 nm using a CoolSnap camera (CoolSnap HQ, Visiiton, Puchheim, Germany). [Ca2+]i was calculated from the 340/380 nm fluorescence ratio after background subtraction. The formula used to calculate [Ca2+]i was [Ca2+]i = Kd × (R – Rmin) / (Rmax – R) × (F2/Fb2), where R is the observed fluorescence ratio. The values Rmax and Rmin (maximum and minimum
ratios) and the constant $S_{f2}/S_{b2}$ (fluorescence of free and $Ca^{2+}$-bound Fura-2 at 380 nm) were calculated using 1 $\mu$M ionomycin (Calbiochem, Merck KGaA, Darmstadt, Germany), 5 $\mu$M nigericin, 10 $\mu$M monensin (Sigma-Aldrich, Merck KGaA), and 5 mM EGTA to equilibrate intracellular and extracellular $Ca^{2+}$ in intact Fura-2–loaded Paneth cells. The dissociation constant for the Fura-2–$Ca^{2+}$ complex was taken as 224 nmol/L.

Control of experiments, imaging acquisition, and data analysis were done with the software package Meta-Fluor (Molecular Devices, Biberach, Germany).

**Materials and Statistical Analysis**

All compounds used were of highest available grade of purity and were bought from Sigma-Aldrich (Merck KGaA, Darmstadt, Germany), unless indicated otherwise. Data are shown as individual traces/representative images and/or as summaries with mean values $\pm$ standard error of the mean, with the respective number of experiments given in each figure legend. For statistical analysis, paired or unpaired Student’s t-test, analysis of variance (post hoc Bonferroni), or Kruskal-Wallis test (H-Test) were used as appropriate. A $P$-value of $< .05$ was accepted as a statistically significant difference.

**Results**

**Accumulation of Lysozyme in Paneth Cells of Mice Lacking Expression of Tmem16a or Tmem16f**

In previous studies, we detected the expression of Tmem16a and Tmem16f in mouse intestinal epithelial cells. Here, we analyzed the expression of Tmem16a and Tmem16f in the jejunal crypt base of wild-type (WT), TMEM16a$^{-/-}$, and Tmem16f$^{-/-}$ mice. Tmem16a was detected at the apical pole, whereas expression of Tmem16f was located predominantly at the basolateral side of crypt epithelial cells (Figure 1A, Figure A1). Notably, Tmem16a is also located in intestinal smooth muscle cells. We detected a pronounced accumulation of lysozyme in jejunal Paneth cells of TMEM16a$^{-/-}$ and Tmem16f$^{-/-}$ mice when compared with WT animals (Figure 1B and C). These results suggested a Paneth cells secretory defect in the absence of TMEM16a or Tmem16f. Paneth cells at the crypt base can be easily identified by their abundant eosinophilic granules. When analyzing small intestinal Paneth cells, we detected an enhanced size of Paneth cells in Tmem16f$^{-/-}$ mice (Figure 2A and B). Moreover, the number of Paneth cells per crypt and the number of granules within each Paneth cell were enhanced in TMEM16a$^{-/-}$ and Tmem16f$^{-/-}$ mice when compared with WT (Figure 2A and C).

**Accumulation of Intestinal Mucus in Tmem16a$^{-/-}$ and Tmem16f$^{-/-}$ Mice**

Previous studies demonstrated airway goblet cell metaplasia, an accumulation of mucus in airway and intestinal goblet cells of mice with tissue-specific knockout of Tmem16a and Tmem16f. A more detailed analysis of mucus using periodic acid–Schiff staining of duodenum,
jejunum, and ileum performed in the present study confirmed previous results and demonstrated enhanced mucus in small intestine of TMEM16a−/− and Tmem16f−/− mice (Figure 3, Figure A2). The data suggest that lack of either Tmem16a or Tmem16f causes a broad secretory defect in secretory cells, including Paneth cells.

**Figure 2.** Number and size of Paneth cells are enhanced in Tmem16a−/− and Tmem16f−/− mice. (A) HE staining of crypt bases of duodenum, jejunum, and ileum of WT, T16a−/−, and T16f−/− mice. Bar = 20 μm. (B) Quantitative analysis of Paneth cell sizes in duodenum, jejunum, and ileum. (C, D) Analysis of Paneth cell number and number of granules within Paneth cells in duodenum, jejunum, and ileum of WT, T16a−/−, and T16f−/− mice. Mean ± standard error of the mean (number of animals and crypts analyzed). #significant difference from WT (P < .05, analysis of variance). WT, wild type.

ATP-Induced but not CCH-Induced Release of Granules From Paneth Cells is Compromised by Knockout of Tmem16a

Accumulation of lysozyme in the knockout animals prompted us to examine in more detail the potential secretory defect in Paneth cells. To that end, we generated
jejunal organoids by growing freshly isolated jejunal crypts in a three-dimensional matrigel. These organoids demonstrate features of naïve crypts with enterocytes, goblet cells, and lysozyme-expressing Paneth cells (Figure 4A). Granule-filled Paneth cells could be easily detected by life differential interference contrast microscopy (Figure 4B). In online experiments, organoids were exposed to the secretagogue CCH and to the purinergic ligand ATP. In WT organoids, CCH induced a sudden and complete release of granules from Paneth cells (Figure 4B, WT_CCH.mp4). In contrast, stimulation with ATP induced a slower and only partial release of granules in the apical compartment of Paneth cells (Figure 4B, WT_ATP.mp4). Notably, apical application of ATP by injection into the lumen of organoids caused a similar apical release of granules (WT_ATP.apical injection.mp4). ATP-induced Paneth cell secretion was...
strongly Tmem16a dependent, as almost no ATP-induced secretion was observed in organoids of Tmem16a-knockout mice (Figure 4B and C, KO_ATP.mp4). In contrast, CCH caused Paneth cell secretion even in the absence of Tmem16a, although the release was somewhat delayed (Figure 4B and C, KO_CCH.mp4). These results establish a novel regulation of apical exocytosis of granules in Paneth cells Tmem16a/f.

**Differential Requirement of Tmem16a and Tmem16f for ATP- and CCH-Induced Release of Granules**

We further examined how granular release by ATP and CCH depend on both Tmem16a and Tmem16f. In the absence of Tmem16a, ATP-induced release was strongly attenuated, whereas knockout of Tmem16f did not affect Paneth cell secretion (Figure 5A and B). These results correspond to the primarily apical expression of purinergic receptors28,29 and the apical localization to Tmem16a (Figure 1A). In contrast, muscarinic M3 receptors and Tmem16f are both located at the basolateral pole of Paneth cells30 (Figure 1A). Granular release induced by muscarinic stimulation (CCH) was hardly affected by knockout of apically located Tmem16f but was significantly reduced in the absence of Tmem16f (Figure 5C). Notably, niclosamide, a potent inhibitor of both Tmem16a and Tmem16f,20,31 additionally inhibited granular release by ATP or CCH in Paneth cell lacking Tmem16f. The results demonstrate the importance of both Tmem16a and Tmem16f for fusion of granules in Paneth cells and release of their content into the crypt lumen.

**Enhanced Bacterial Content and Attenuated Programmed Cell Death in Jejunum and Ileum of Tmem16a^-/- and Tmem16f^-/- Mice**

Granules of Paneth cells contain antimicrobial peptides, cytokines, and other factors that control proliferation or epithelial cell death.3,5,32,33 We therefore analyzed the presence of Gram-positive and Gram-negative bacteria in jejunum and ileum of WT, Tmem16a^-/-, and Tmem16f^-/- mice. The number of bacteria was enhanced in the ileum of...
Tmem16a<sup>−/−</sup> and Tmem16f<sup>−/−</sup> mice and in the jejunum of Tmem16f<sup>−/−</sup> mice, suggesting a reduced antimicrobial activity in the absence of Tmem16 proteins (Figure 7). We also compared regulated cell death of intestinal epithelial cells in jejunum of WT and Tmem16 knockout animals. Using TUNEL assays, we found a largely reduced number of cell death in both Tmem16a<sup>−/−</sup> and Tmem16f<sup>−/−</sup> mice. Taken together, the present results unmasked a crucial role of Tmem16a and Tmem16f for Paneth cell secretion. Tmem16a/f maintain secretory intracellular Ca<sup>2+</sup> signals highly relevant for exocytosis of granules and proper Paneth cell function. A diversity of small molecules and natural compounds exist that either activate or inhibit Tmem16a and Tmem16f. The present findings may therefore provide the basis for a novel anti-inflammatory therapy for intestinal diseases and may improve our understanding of the molecular mechanism of some of the currently available drugs.

**Discussion**

**Exocytosis is Controlled by Tmem16a and Tmem16f**

Paneth cells are abundant in the crypt base of small intestine and are occasionally found in the proximal colon. A series of studies have elucidated secreted products that are released by Paneth cells. The most well-known granule contents are the antimicrobial peptides such as lysozyme and α-defensins, phospholipase A2, cytokines (interleukin 17A and tumor necrosis factor-α) and proteases such as metalloproteinase. Granules are exocytosed on contact to luminal LPS, which is a relatively slow process, suggesting a constitutive fusion of single granules with the apical membrane. However, LPS can induce ATP release, and we therefore propose that LPS-induced ATP release triggers regulated (sometimes called constitutive) exocytosis by increasing Ca<sup>2+</sup> in the apical Paneth cell compartment. In contrast, sudden massive compound exocytosis can be triggered by stimulation of basolateral M3 receptors similar to compound exocytosis of mucus in goblet cells. Compound exocytosis is characterized by prefusion of granules, which provides a mechanism where deeper lying granules can readily release their content without having to be transported to the apical cell membrane. The present study shows that both Tmem16a and Tmem16f are required for proper exocytosis. We propose that ATP could induce the release of exocytic granules from Paneth cells or goblet cells via so-called regulated exocytosis, which requires...
apical TMEM16A (c.f. Videos WT_ATP.mp4 and KO_ATP.mp4). TMEM16A is inhibited by Ani9 and by AO1, which also inhibits regulated exocytosis. In contrast, CCH possibly induces the release of Paneth cell granules by compound exocytosis, which requires the function of basolateral TMEM16F18 (c.f. Videos WT_CCH.mp4 and KO_CCH.mp4). In contrast to TMEM16A, TMEM16F is not inhibited by AO1 or Ani9. AO1 and Ani9 inhibited not only TMEM16A and regulated exocytosis but also inhibited ATP-induced Ca$^{2+}$ signaling.16,18,41 Taken together, Tmem16a, being located in the apical membrane, appears to be more relevant for regulated exocytosis, whereas Tmem16f located at the basolateral pole could be important for compound exocytosis.

Intracellular Ca$^{2+}$ Signals are Shaped by Tmem16 Proteins

Both Tmem16a and Tmem16f may control mucus release in intestinal and airway goblet cells by different mechanisms.10,20,23 In large intestine, we and others found expression of TMEM16A predominantly, but not exclusively, in basolateral (particularly lateral) membranes of colonic crypt cells.18,27,42 It was shown that Tmem16a tethers the ER to the apical membrane. This leads to efficient receptor-mediated Ca$^{2+}$ store release and Ca$^{2+}$-dependent fusion of vesicles/granules with the apical membrane.16,17,43,44 In contrast, Tmem16f is a phospholipid scramblase and an ion channel and could have 2 functions: It might promote membrane fusion to form giant granules, which are known to be formed during compound exocytosis. For example, membrane fusion by Tmem16f was also observed for SARS-CoV-2-induced syncytia formation.45–47 However, Tmem16f also allows permeation of Ca$^{2+}$ and may thereby support ATP-induced increase in intracellular Ca$^{2+}$ as shown in the present study and in previous studies.48

Consequences of Loss of Function of Tmem16a and Tmem16f: Possible Therapeutic Targets in Intestinal Disease

Tmem16a is expressed in numerous tissues, preferentially in epithelial cells. Thus, a loss of function has numerous consequences, including compromised exocytosis, lower expression of proteins in the plasma membrane, and attenuated secretion of mucus and electrolytes.18,20,23,49

In contrast, Tmem16f is highly expressed in macrophages, B-lymphocytes, platelets, and osteoblasts, and therefore,
loss of function causes immune defense, hemostasis, and bone mineralization.\textsuperscript{50–52}

In our intestinal epithelial-specific Tmem16a/f knockout mice, we found an enhanced bacterial content in the intestine, which is related to the attenuated Paneth cell exocytosis and lower release of antimicrobial compounds (Figure 7). We also detected a reduced rate of apoptosis/necroptosis.\textsuperscript{33} Notably, Paneth cells also express and release trophic factors, such as epidermal growth factor and mediators of the Wnt and Notch signaling pathway. These factors regulate intestinal stem cell homeostasis, secretory cell differentiation, and death of aged cells.\textsuperscript{3,32,33} In this context, it is interesting to note that TMEM16F is required for activation of the metalloproteinases ADAM10 and ADAM17.\textsuperscript{53,54} ADAM10 is expressed throughout the intestinal epithelium where it induces shedding of epithelial growth hormone receptors.\textsuperscript{55}

Intestinal inflammatory diseases such as Crohn’s disease, necrotizing enterocolitis, and intestinal microbiota dysbiosis have been related to abnormal Paneth cell physiology.\textsuperscript{3,56} Along this line, we reported the first 2 patients with a loss-of-function mutation in TMEM16A, which suffered from recurrent episodes of hemorrhagic diarrhea and necrotizing enterocolitis.\textsuperscript{57} Meanwhile, many small molecules and numerous natural or herbal compounds have been identified that either inhibit or activate Tmem16a and Tmem16f.\textsuperscript{21,22,31,34,37,58,59} Some of these compounds may turn out to be useful therapeutics in inflammatory bowel disease, intestinal allergies, or abnormal colonization of the gut. Activators of TMEM16A such as the compound ETX001/ETD001 are currently under evaluation for the treatment of cystic fibrosis (CF; \url{https://www.clinicaltrials.gov/}). It is assumed to restore an apical Cl\textsuperscript{−} conductance in airways of CF patients, which lack functional cystic fibrosis conductance regulator Cl\textsuperscript{−} channels. Our team, however, does not support the use of TMEM16A activators in CF airways, as we have shown that activation of TMEM16A leads to an increase in mucus production and mucus secretion\textsuperscript{20,21} and present study. However, activation of TMEM16A and consecutive release from Paneth cells and secretion of protective intestinal mucus might be well beneficial to patients suffering from various forms of inflammatory bowel disease.

**Study Approval**

All animal experiments complied with the ARRIVE guidelines and were carried out in accordance with the UK Animals (Scientific Procedures) Act 1986 and associated guidelines, as well as EU Directive 2010/63/EU for animal experiments. The study was approved by the local Ethics Committee of the Government of Unterfranken (RUF-55.2.2-2532-2-677-19; approved on August 24, 2018), and our investigations were carried out in accordance with the Guide for the Care and authorities at the University of Kiel (project agreement no #1130).

**Supplementary Materials**

Material associated with this article can be found in the online version at \url{https://doi.org/10.1016/j.gastha.2022.08.002}.

**References**


