Drug Inhibition of SARS-CoV-2 Replication in Human Pluripotent Stem Cell–Derived Intestinal Organoids

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SUMMARY

Human pluripotent stem cell–derived intestinal organoids serve as an indefinite resource for organ-specific drug testing. SARS-CoV-2 infected and replicated within different cell types of the organoids, which was effectively inhibited by remdesivir and EK1 but not by famotidine.

BACKGROUND AND AIMS: The COVID-19 pandemic has spread worldwide and poses a severe health risk. While most patients present mild symptoms, descending pneumonia can lead to severe respiratory insufficiency. Up to 50% of patients show gastrointestinal symptoms like diarrhea or nausea, intriguingly associating with prolonged symptoms and increased severity. Thus, models to understand and validate drug efficiency in the gut of COVID-19 patients are of urgent need.

METHODS: Human intestinal organoids derived from pluripotent stem cells (PSC-HIOs) have led, due to their complexity in mimicking human intestinal architecture, to an unprecedented number of successful disease models including gastrointestinal infections. Here, we employed PSC-HIOs to dissect SARS-CoV-2 pathogenesis and its inhibition by remdesivir, one of the leading drugs investigated for treatment of COVID-19.

RESULTS: Immunostaining for viral entry receptor ACE2 and SARS-CoV-2 spike protein priming protease TMPRSS2 showed broad expression in the gastrointestinal tract with highest levels in the intestine, the latter faithfully recapitulated by PSC-HIOs. Organoids could be readily infected with SARS-CoV-2 followed by viral spread across entire PSC-HIOs, subsequently leading to organoid deterioration. However, SARS-CoV-2 spared goblet cells lacking ACE2 expression. Importantly, we challenged PSC-HIOs for drug testing capacity. Specifically, remdesivir effectively inhibited SARS-CoV-2 infection dose-dependently at low micromolar concentration and rescued PSC-HIO morphology.

CONCLUSIONS: Thus, PSC-HIOs are a valuable tool to study SARS-CoV-2 infection and to identify and validate drugs especially with potential action in the gut. (Cell Mol Gastroenterol Hepatol 2021;11:935–948; https://doi.org/10.1016/j.jcmgh.2020.11.003)

Keywords: SARS-CoV-2; COVID-19; Intestinal Organoids; Remdesivir; Famotidine.
Results

**ACE2 and TMPRSS2 Are Expressed in the Gastrointestinal Tract and in PSC-HIOs**

First, we studied expression of ACE2 and TMPRSS2 in organs of the gastrointestinal tract. Expression of both SARS-CoV-2 entry factors was most prominent in the epithelial lining of duodenum, gallbladder, and colon as compared with Caco-2 cells (Figure 1A–C). Duodenal cells showed apical ACE2 expression in the glyocalyx and a strong cytoplasmic TMPRSS2 signal (Figure 1A and B). Cells of gallbladder and colon presented a strong luminal expression of both proteins while sections of gastric mucosa and esophagus only revealed weak expression (Figure 1C). These findings suggest that various cell types in gastrointestinal tissues are SARS-CoV-2 target cells, though with different expression intensities of the entry factors. Next, we analyzed ACE2 and TMPRSS2 expression in in vitro differentiated PSC-HIOs (Figure 1D–H). Both proteins were readily expressed in virtually all intestinal cells present in PSC-HIOs (Figure 1E–H) mimicking expression patterns in human tissue and Caco-2 cells (Figure 1A and B). Moreover, co-staining for distinct intestinal cell type markers revealed that ACE2 is expressed on chromogranin A (CHGA)-positive enteroendocrine and lysozyme (LYZ)-positive Paneth cells (Figure 1F and G), while mucin 2 (MUC2)-positive goblet cells appeared virtually negative (Figure 1H). Thus, PSC-HIOs express cell-specifically SARS-CoV-2 entry factors and might be susceptible to infection.

**PSC-HIOs Support SARS-CoV-2 Infection and Replication**

To determine susceptibility to in vitro infection, PSC-HIOs were exposed to SARS-CoV-2, and expression of viral spike (S) and nucleocapsid (N) protein together with the epithelial marker E-cadherin (Ecad) or respective labels for more specialized cell types (CHGA, LYZ, MUC2) were analyzed. Upon SARS-CoV-2 infection of PSC-HIOs, S protein was detected in 10% of the cells after 24 hours, which increased to 57% at 48 hours, suggesting viral replication and spreading infection (Figure 2A and B). Costaining for N protein confirmed this finding (Figure 2C). Notably, infected cells are positive for cleaved caspase 3 (CASP3) indicating initiated apoptosis in SARS-CoV-2-positive intestinal cells.
Discussion

We report that in vitro differentiated PSC-HIOs express ACE2 and TMPRSS2 and support productive infection with SARS-CoV-2 in a cell type-specific manner, launching a valuable pathomechanistic model for SARS-CoV-2 infection of the gut. Notably, remdesivir and EK1 but not famotidine block SARS-CoV-2 in PSC-HIOs, indicating suitability to treat gastrointestinal COVID-19 and validating PSC-HIOs as a putative screening platform to test drugs that might ameliorate gastrointestinal symptoms in the clinic.

A study by Lamers et al. recently demonstrated SARS-CoV-2 infection of enterocytes in small intestinal organoids from primary gut epithelial cells. Similarly, recent studies show infection of and replication in epithelial cells of human colon-derived intestinal organoids as well as human and bat small intestinal organoids. We extended these studies by specifically showing infection of specialized intestinal cells (e.g., enteroendocrine and Paneth cells) in addition to enterocytes, suggesting that SARS-CoV-2 infection may not only disrupt absorption and transport of metabolites, but also hamper hormone secretion and local immune defense. Similarly, infection of Paneth cells may support further disturbance of host defense by impeding secretion of antiviral and antimicrobial proteins into the gut. However, qPCR analysis of cell type–specific markers did not reveal conclusive data, potentially influenced by the sampling time point. Especially considering the recently highlighted relevance of mucosal SARS-CoV-2 defense to fight COVID-19, our findings of cell type–specific mucosal

Drug Testing in PSC-HIOs

We next analyzed the effect of remdesivir on SARS-CoV-2 infection of intestinal cells. On Caco-2 cells, remdesivir inhibited viral infection in a dose-dependent manner with a half-maximal inhibitory concentration (IC$_{50}$) of 46 nM (Figure 4A). Cell viability remained unaffected up to concentrations of 25 μM of the drug and was only marginally affected at 125 μM, resulting in a selectivity index value for remdesivir in colorectal Caco-2 cells of >2,717 (Figure 4A and B). Notably, remdesivir also decreased SARS-CoV-2 infection of PSC-HIOs by 86% at a concentration of 500 nM and almost completely abolished infection at 5 μM (Figure 4C and D). Accordingly, viral copies as measured via qPCR for CoV-N were strongly diminished at a dosage of 5 μM (Figure 4E). We also determined IC$_{50}$ values by stepwise dose escalation of remdesivir in SARS-CoV-2–infected PSC-HIOs by flow cytometry analysis of N protein expression (Figure 4F). In contrast to the rather low IC$_{50}$ value of 46 nM in monolayer cultured Caco-2 cells, the IC$_{50}$ value in PSC-HIOs was ~70-fold higher with 3.2 μM, indicating the importance of performing drug tests in a 3D culture format to avoid overestimations (Figure 4A and F). With an IC$_{50}$ of 3.2 μM and a 50% cytotoxic concentration exceeding 125 μM of remdesivir on PSC-HIOs, however, the selectivity index is at least higher than 39, indicating good tolerability (Figure 4G). Tissue culture infection dose 50 (TCID$_{50}$) determination of the supernatants derived from drug-treated infected PSC-HIOs confirmed a dose-dependent inhibition of SARS-CoV-2 replication (Figure 4H).

The over-the-counter histamine-2 receptor antagonist famotidine is a putative therapeutic agent for COVID-19. Thus, we tested famotidine concentrations of up to 2.5 mM but did not observe notable reduction of SARS-CoV-2 infection of Caco-2 cells (Figure 5A). In contrast to remdesivir, famotidine compromised Caco-2 cell viability starting from 5 mM upward (Figure 5B). Accordingly, PSC-HIOs did not show reduced infection rates after famotidine treatment, as assessed by immunostaining for viral S protein and CoV-N qPCR (Figure 5C, D, and F). Finally, we evaluated the antiviral activity of EK1, a recently described peptidic pan-coronavirus fusion inhibitor, in PSC-HIOs. In a previous study, we found EK1 to inhibit SARS-CoV-2 infection of Caco-2 with an IC$_{50}$ of 303.5 nM, while not causing any cytotoxic effects up to 10 μM concentration (selectivity index > 32.9). Accordingly, in PSC-HIOs the presence of 10 μM EK1 during infection decreased the number of S-positive cells 48 hours after infection by 38% (Figure 5E and F). Taken together, SARS-CoV-2 productively infects most gastrointestinal cell types, with the notable exception of goblet cells, and remdesivir and the peptidic pan-coronavirus fusion inhibitor EK1 but not famotidine block SARS-CoV-2 infection of intestinal organoids.
Figure 2. PSC-HIOs support SARS-CoV-2 infection and replication. (A) PSC-HIOs were infected with SARS-CoV-2 and stained for viral S protein (red) and E-cadherin (Ecad) (green) at day 1 and day 2 postinfection. Nuclei are stained with DAPI in blue. Scale bar = 50 μM. (B) Infected cells of PSC-HIOs were manually counted according to presence of viral S protein compared with the total cell number (n = 3 independent experiments, at least 3 organoids counted per experiment). Infection was further confirmed by S (red) and N (green) protein co-staining. Scale bar = 50 μM. (D) Infected cells of PSC-HIOs 2 days postinfection were stained for viral S protein (red) and CASP3 (green). Nuclei are stained with DAPI in blue. Scale bar = 50 μM. (E) Cleaved caspase 3 (CASP3)-positive cells were quantified by flow cytometry (n = 3). (F) Relative SARS-CoV-2 viral RNA copies in PSC-HIOs at different time points were quantified using primers amplifying CoV-N in RT-qPCR. For normalization, Ct values obtained for amplification of GAPDH from the same cultures were subtracted from the Ct value obtained for CoV-N (n = 2). (G) Viral genome copy number in extracellular matrix (matrigel) of infected PSC-HIOs was assessed by RT-qPCR targeting CoV-N and -ORF1b-nsp14 (n = 2). All error bars represent SEM.

Figure 1. (See previous page). ACE2 and TMPRSS2 are expressed in the gastrointestinal tract and in PSC-HIOs. (A) Paraffin-embedded sections of indicated gastrointestinal tissues and embedded Caco-2 cells were stained for ACE2 and TMPRSS2 protein. Strong luminal expression is found in epithelia of duodenum, colon, and gallbladder, with weaker expression in the stomach and the mucosal lining of the esophagus. Representative images are shown. Scale bars = 20 μm. (B) Duodenum biopsy was stained for ACE2 and TMPRSS2 using primary antibodies and fluorophore-conjugated secondary antibodies and imaged by fluorescence microscopy. Scale bar = 50 μM. (C) ACE2 and TMPRSS2 expression from panel A was graded according to signal intensity (relative to the positive control, Caco-2 cells). (D) Microscopy of PSC-HIOs. Scale bar = 50 μM. (E) PSC-HIOs were stained for ACE2 (left panels) and TMPRSS2 (right panels). Nuclei are stained with DAPI in blue. (F–H) PSC-HIOs were costained for ACE2 and (F) enteroendocrine cell marker CHGA, (G) Paneth cell marker LYZ, or (H) goblet cell marker MUC2 showing coexpression with CHGA and LYZ but not with MUC2. Nuclei are stained with DAPI in blue. Arrows indicate co-expression. Left panels show representative images, right panels magnifications.
SARS-CoV-2 infection add information about SARS-CoV-2 tropism. Our infection pattern of SARS-CoV-2 sparing goblet cells contrasts with lung findings from deceased COVID-19 patients, in which MUC5AC-positive goblet cells as well as MUC5B-positive club-like cells have been infected.34 These tissue-specific differences should be subject to further investigation.

As COVID-19 patients suffering from gastrointestinal symptoms show prolonged disease duration and severity of disease, cells such as enteroendocrine cells, Paneth cells, and goblet cells are particularly relevant for local immune defense; thus, drugs that suppress intestinal SARS-CoV-2 infection are of high interest. Remdesivir, developed to target Ebola virus, provides the first antiviral attempt to treat COVID-19, although clinical success appears heterogeneous and benefiting subpopulations remain to be defined.25,35 Similarly, effectivity of this agent across distinct organs is completely unclear. We here show that remdesivir inhibits SARS-CoV-2 infection of and replication in PSC-HIOs, highlighting its possible use for treatment of gastrointestinal infection occurring simultaneously with or independently of respiratory manifestation of COVID-19. In addition, head-to-head comparison of IC50 values determined in a 3D vs 2-dimensional culture system underpins the relevance of screening system, as 2-dimensional cultures are likely overestimating efficiency of a given drug.

In contrast, famotidine, proposed to inhibit proteins essential for viral replication,26 did not affect SARS-CoV-2 infection and viral spread in a colon cell line and PSC-HIOs. However, our current treatment regime does not allow to determine long-term effects, which may play a role during clinical deterioration. This would be in line with low binding affinity of famotidine to SARS-CoV-2 proteases28; however, its effect on host-specific immune response as well as tissue-specific efficiency might be of relevance to ultimately judge its clinical value, particularly to treat life-limiting lung infection. In line, histamine-2 receptor antagonists are able to modulate histamine effector pathways by receptor binding, thus directing immunomodulatory effects.36 Specifically, they can boost the innate immune function and revert negative effects of histamine on production and release of cytokines such as tumor necrosis factor alpha and interferon alpha, possibly supporting immune response during viral infection.37,38 Accordingly, a recent case series suggested efficiency to treat mild-to-moderate COVID-19 with pulmonary parameters as a readout.29 Thus, the current clinical study landscape remains in support of famotidine, although our in vitro data appear contrary and warrant further investigation in randomized controlled trials, best with taking gastrointestinal facets of COVID-19 into account.

Conclusions

Overall, our data demonstrate that SARS-CoV-2 can infect all investigated cell types of the gut, with the exception of goblet cells, and can perturb intestinal integrity, which might lead to diarrhea among many other gastrointestinal symptoms frequently reported by COVID-19 patients. Most importantly, we tested and validated our culture platform as an indefinite resource to organ-specific drug testing, as shown by proof-of-concept studies with remdesivir and EK1, emphasizing the relevance of the system as a potential tool for drug screening. Treatment with remdesivir during in vitro infection of intestinal organoids with SARS-CoV-2 inhibited viral replication. Hence, clinical treatment with this agent might prevent SARS-CoV-2-induced gastrointestinal damage and soothe gastrointestinal symptoms.

Materials And Methods

Drugs

Remdesivir was obtained from Selleck Chemicals (#S8932; Houston, TX), famotidine was obtained from Sigma-Aldrich (#F6889-1G; St. Louis, MO), and EK1 was synthesized by the Core Facility Functional Peptidomics, Ulm University Medical Center.

Cell Culture

Caco-2 cells were grown in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% heat-inactivated fetal calf serum (FCS), 100 units/mL penicillin, 100 μg/mL streptomycin (Sigma-Aldrich), 2 mM L-glutamine, 1 mM sodium pyruvate, and NEAA (#M7145; Sigma-Aldrich). Vero E6 cells were grown in the same medium but with 2.5% FCS. TMPRSS2-expressing Vero E6 cells (kindly provided by the National Institute for Biological Standards and Control, Ridge, United Kingdom; #100978) were cultivated in DMEM supplemented with 10% FCS, 2 mM L-glutamine, 100–units/mL penicillin, 100 mg/mL streptomycin, and 1 mg/mL geneticin. Cells were grown at 37°C in a 5% CO2 humidified incubator.

Stem Cell Culture and Intestinal Differentiation

Human embryonic stem cell line HUES8 (Harvard University, Cambridge, MA) was used with permission from the Robert Koch Institute according to the “79. Genehmigung

Figure 3. (See previous page). SARS-CoV-2 infects enteroendocrine and Paneth cells but not goblet cells. (A–F) Confocal microscopy (left column; A, C, E) and flow cytometric analysis (right column; B, D, F) of SARS-CoV-2 infected cells costained for S protein (red) and markers for goblet (green, MUC2), enteroendocrine (green, CHGA), and Paneth cells (green, LY2). Scale bar = 20 μm. (G) Flow cytometry–based quantifications of the frequency of indicated cell types in intestinal organoids are shown. “Other” cells comprise mostly enterocytes and few stem cells.17,18 (H) Flow cytometric quantification of infected (S-positive) PSC-HIO cell types compared with total cell number (2 independent experiments with 2–3 biological replicates per experiment are shown; error bar represents SEM). (I) RT-qPCR quantification of enteroendocrine, Paneth cell–, goblet cell–, and stem cell–associated transcripts shown as fold change following infection by SARS-CoV-2 (compared with uninfected cells; 2 independent experiments with 2 biological replicates per experiments).
nach dem Stammzellgesetz, AZ 3.04.02/0084.” Cells were cultured on human embryonic stem cell matrigel (Corning, Corning, NY) in mTeSR1 medium (STEMCELL Technologies, Vancouver, Canada) at 5% CO₂, 5% O₂, and 37°C. Medium was changed daily and cells were split with TrypLE Express (Invitrogen, Carlsbad, CA). For differentiation, 300,000 cells
per well were seeded in 24-well plates coated with growth factor-reduced matrigel (Corning) in mTeSR1 with 10-μM Y-27632 (STEMCELL Technologies). The next day, differentiation was started at 80%-90% confluency, as reported previously.13,18,39–41

**Virus Strains and Virus Propagation**

Viral isolate BetaCoV/Netherlands/01/NL/2020 (#010V-03903) was obtained from the European Virus Archive global and propagated on Vero E6 cells. To this end, 70%-90% confluent cells in 75 cm² cell culture flasks were inoculated with 100-μL SARS-CoV-2 isolate in 3.5-mL serum-free medium containing 1-μg/mL trypsin. Cells were incubated for 2 hours at 37°C, before adding 20-mL medium containing 15-mM HEPES. Cells were incubated at 37°C and supernatant harvested when a strong cytopathic effect was visible. Supernatants were centrifuged for 5 minutes at 1,000 g to remove cellular debris, and then aliquoted and stored at -80°C as virus stocks. Infectious virus titer was determined as plaque-forming units on Vero E6 cells, which was used to calculate multiplicity of infection.

**In Vitro Infection and Cell-Based SARS-CoV-2 Immunodetection Assay**

To determine SARS-CoV-2 infection, 30,000 Caco-2 cells were seeded in 96-well plates. The next day, the compound of interest was added and the cells were infected with a multiplicity of infection of 0.001 in a total volume of 180 μL. Two days later immunodetection assay was performed as previously described32; cells were fixed in 4% paraformaldehyde (PFA), permeabilized using 0.1% Triton-X, and stained with 1:5000 diluted anti-SARS-CoV-2 S protein antibody 1A9 (Biozol GTX-GTX632604) in antibody buffer (10% FCS and 0.3% Tween 20 in phosphate-buffered saline [PBS]) for 1 hour at 37°C. After 3 washes, the secondary horseradish peroxidase–conjugated antibody (#A16066; Thermo Fisher Scientific, Waltham, MA) (1:15,000) was incubated for 1 hour at 37°C. Following 4 times of washing, the TMB peroxidase substrate (#52-00-04; Medac, North Augusta, SC) was added for 5 minutes and the reaction stopped using 0.5 M H2SO4. The optical density was recorded at 450–620 nm using the Asys Expert 96 UV microplate reader (Biochrom, Cambridge, United Kingdom). Values were corrected for the background signal derived from uninfected cells and untreated controls were set to 100% infection.

**In Vitro Cell Viability Assay**

The effect of analyzed compounds on the metabolic activity of Caco-2 cells was analyzed using the CellTiter-Glo Luminescent Cell Viability Assay (#G7571; Promega, Madison, WI) as previously described.

**Infection of Organoids and Drug Testing**

To prepare in vitro differentiated organoids for infection, matrigel was dissolved in Collagenase/Dispase (Roche, Basel, Switzerland) for 2 hours at 37°C and stopped by cold neutralization solution (DMEM, 1% bovine serum albumin, and 1% penicillin-streptomycin). Organoids were transferred into 1.5-mL tubes and infected in 350-μL virus inoculum containing 3 × 10⁶ plaque-forming units SARS-CoV-2 with or without drugs for 1 hour at 37°C. Organoids were then resuspended in 35-μL cold growth factor–reduced matrigel to generate cell-matrigel domes in 48-well plates. After 10 minutes at 37°C, intestinal growth medium (DMEM F12 [Gibco, Gaithersburg, MD], 1× B27 supplement [Thermo Fisher Scientific], 2-mM L-glutamine, 1% penicillin-streptomycin, 40 mM HEPES [Sigma-Aldrich], 3 μM CHIR99021, 200 mM LDN-193189 [Sigma-Aldrich], 100 ng/mL hEGF [Novoprotein, Summit, NJ], and 10 μM Y-27632 [STEMCELL Technologies]) was added and organoids were incubated at 37°C.

**Flow Cytometry Analysis of Organoids**

After infection and drug treatment, matrigel was dissolved in Collagenase/Dispase (Roche) for 2 hours at 37°C and stopped by cold neutralisation solution (DMEM, 1% bovine serum albumin, and 1% penicillin-streptomycin). Single cells were generated by incubation with accutase.
Figure 5. EK1 but not famotidine inhibit infection of PSC-HIOs with SARS-CoV-2. (A) Caco-2 cells were infected with SARS-CoV-2 and treated with famotidine at indicated concentrations. Two days later, infection was quantified by an enzyme-based immunodetection assay against viral S protein. Values shown represent mean values derived from triplicate infections ± SEM from one representative experiment. Experiment was performed 2 times. (B) Effect of famotidine on metabolic activity of Caco-2 cells was assessed by CellTiter-Glo assay. Values shown represent mean values derived from triplicates ± SEM from one representative experiment. (C–E) PSC-HIOs were infected with SARS-CoV-2 and treated with (C) famotidine or (E) EK1. Two days later, infection was analyzed by staining for viral S protein (red). Ecad (green) was additionally stained to visualize cells. Nuclei are stained with DAPI in blue. Scale bar = 50 μm. (D) SARS-CoV-2 N was quantified in cellular RNA isolates of PSC-HIOs treated with either distinct dosages of famotidine and normalized to GAPDH RNA (n = 1, values represent means ± SEM of technical duplicates). (F) Images from panels C and E were quantified according to the presence of viral S protein compared with total cell number (n = 2–3 independent experiments, at least 3 organoids counted per experiment; values represent mean ± SEM).
(STEMCELL Technologies) for 30 minutes at 37°C. Cells were fixed in 4% PFA for 1 hour. For flow cytometry analysis, cells were blocked in 5% normal donkey serum in 0.1% Triton-X in PBS for 30 minutes on ice. Cells were incubated with primary antibodies diluted in blocking solution overnight at 4°C. After washing twice with 2% normal donkey serum in 0.1% Triton-X in PBS, cells were incubated with secondary antibodies (Alexa Fluor IgG HþL [Invitrogen]; 1:500) in blocking solution for 1.5 hours on ice in the dark. Cells were washed twice and filtered, and fluorescence was measured using a BD LSR II flow cytometer (BD Biosciences, San Jose, CA).

**Annexin V Viability Assay**

Organoids were treated with drugs for 48 hours. Matrigel was dissolved in Collagenase/Dispase (Roche) for 2 hours at 37°C and stopped by cold neutralization solution (DMEM, 1% bovine serum albumin, and 1% penicillin-streptomycin). Single cells were generated by incubation with accutase (STEMCELL Technologies) for 30 minutes at 37°C. Cells were stained using the FITC Annexin V Apoptosis Detection Kit (BD Biosciences) according to manufacturer’s instructions and analyzed using a BD LSR II Flow cytometer (BD Biosciences).

**Isolation of RNA From Matrigel and Organoids for RT-qPCR**

Viral RNA from matrigel of infected organoids or organoids themselves was isolated using the Qiagen Viral RNA Mini Kit (#52906; Qiagen, Hilden, Germany) or Qiagen RNeasy Plus Mini Kit (#74136; Qiagen), respectively, both as described by the manufacturer. For this purpose, matrigel was digested as described previously, mixed with AVL buffer and incubated at room temperature for 20 minutes before freezing at -20°C. Similarly, matrigel was dissolved and removed and organoids were lysed in 600-μL RLT Plus buffer containing 1% β-mercaptoethanol, vortexed for 30 seconds and then frozen at -20°C until analysis. Organoid lysates were homogenized using QIAshredder (#79656; Qiagen).

**Reverse-Transcription qPCR**

RT-qPCR from matrigel was performed with primer sets targeting N (nucleoprotein) and ORF1b-nsp14 (ORF1b)42 as previously described43 using TaqMan Fast Virus 1-Step Master Mix (#4444436; Thermo Fisher Scientific) and a StepOnePlus Real-Time PCR System (96-well format, fast mode; Thermo Fisher Scientific). Synthetic SARS-CoV-2 RNA (#102024 [Twist Bioscience, San Francisco, CA] or #VR-3276SD [American Type Culture Collection, Manassas, VA]) was used as a standard to obtain viral copy numbers. For RNA isolated from organoids, GAPDH was analyzed as an endogenous control (#4310884E; Applied Biosystems, Foster City, CA) and resulting Ct values subtracted from those obtained for CoV-N reactions for the same samples. All reactions were run in duplicates.

For RT-qPCR of intestinal cell type transcripts, RNA was transcribed into cDNA using the iScript cDNA synthesis Kit (170-8891; Bio-Rad, Hercules, CA). RT-qPCR was performed using Qiagen QuantiTect Primer Assays and SensiMix SYBR No Rox Kit (QT650-02; Bioline, London, United Kingdom) in 96-well format. Expression was normalized to hydroxymethylbilane synthase.

**TCID<sub>50</sub> Endpoint Titration**

To determine the TCID<sub>50</sub>, supernatant samples were taken at day 0 and day 2. Samples were serially diluted and used to inoculate target cells. To this end, 20,000 Vero E6 or TMPRSS2-expressing Vero E6 cells were seeded per well in 96 flat-bottom well plates in 100 μL of medium and incubated over night before 62 μL of fresh medium was added. Next, 18 μL of 5-fold titrated sample was used for inoculation in triplicates. Cells were then incubated for 6 days and monitored for cytopathic effect. TCID<sub>50</sub>/mL was calculated according to Reed and Muench.44

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**Table 1. Primary Antibodies**

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ACE2, angiotensin-converting enzyme 2; CASP3, caspase 3; CHGA, chromogranin A; Ecad, E-cadherin; LYZ, lysozyme; MUC2, mucin 2; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2; TMPRSS2, transmembrane serine protease 2.
Histology of Organoids and Tissue Sections

Sections of human gastrointestinal tissues were provided by the pathology department of Ulm University. Experiments were conducted in accordance with guidelines of the Ethics Committee of the Federal General Medical Council and approved by the Ethics Committee of Ulm University. For histological examination of organoids, they were fixed in 4% PFA over night at 4°C, washed with PBS, and pre-embedded in 2% agarose (Sigma-Aldrich) in PBS. After serial dehydration, intestinal organoids were embedded in paraffin, sectioned at 4 μm, deparaffinized, rehydrated, and subjected to heat-mediated antigen retrieval in tris buffer (pH 9) or citrate buffer (pH 6). Tissue was permeabilized with 0.5% Triton-X for 30 minutes at room temperature and stained overnight with primary antibodies (Table 1) in antibody diluent (ZYTOMED Systems, Berlin, Germany) in a wet chamber at 4°C. After washing with PBS with Tween 20, slides were incubated with secondary antibodies (Alexa Fluor IgG H+L; 1:500) and 500-ng/mL DAPI in Antibody Diluent for 90 minutes in a wet chamber at room temperature. After washing with PBS with Tween 20 and water, slides were mounted with Fluoromount-G (Southern Biotech, Birmingham, AL). Negative controls were performed using IgG controls or irrelevant polyclonal serum (anti-mycobacterium tuberculosis) for polyclonal antibody diluent (ZYTOMED Systems, Berlin, Germany) in a wet chamber at 4°C. After washing with PBS with Tween 20, slides were mounted with Fluoromount-G (Southern Biotech, Birmingham, AL). Negative controls were performed using IgG controls or irrelevant polyclonal serum (anti-mycobacterium tuberculosis) for polyclonal antibodies, respectively. Absence of background staining confirmed specificity of the primary antibodies. Images were acquired with Keyence BZ 9000 microscope (Keyence, Osaka, Japan) and quantified with ImageJ (Version 2.0.0, National Institutes of Health, Bethesda, MD). Laser Scanning confocal images were acquired using the Zeiss LSM710. Z-stacks were deconvoluted and edited in Huygens (Scientific Volume Imaging, Hilversum, the Netherlands) and Fiji (ImageJ).

Statistics

Statistics were performed using GraphPad Prism (GraphPad Software, San Diego, CA). IC50 was determined by nonlinear regression (inhibitor) vs normalized response. For Figure 3G, 4D, and 5D 1-way analysis of variance and Dunnett’s multiple comparison test were performed. Data of Figure 2E were analyzed with a paired t test. *P values <.05 were classified as significant (*P < .05, **P < .01, ***P < .001, ****P < .0001).

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