

Supplemental Information

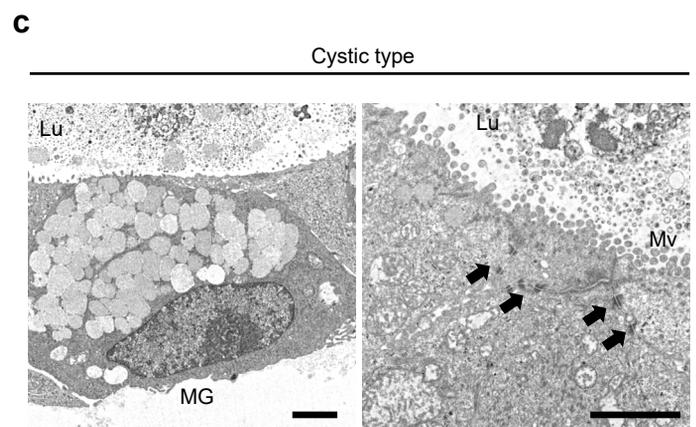
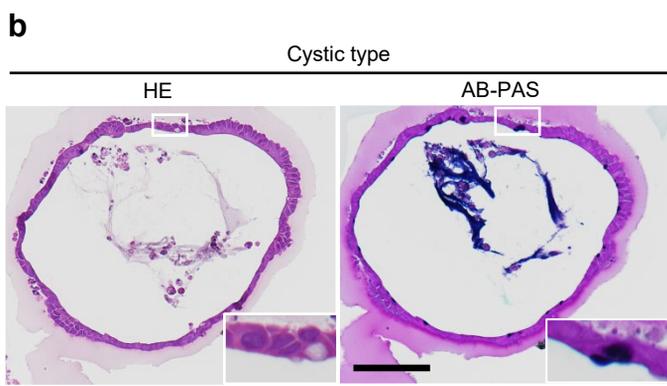
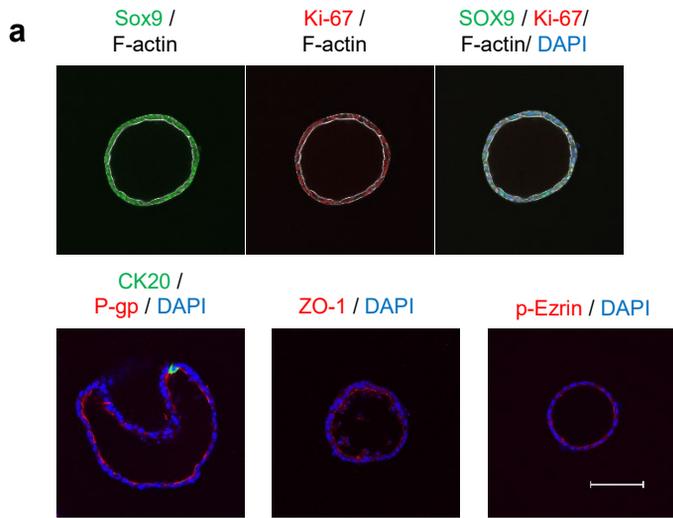
Development of rat duodenal monolayer model with effective barrier function from rat organoids for ADME assay

Authors

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Affiliations

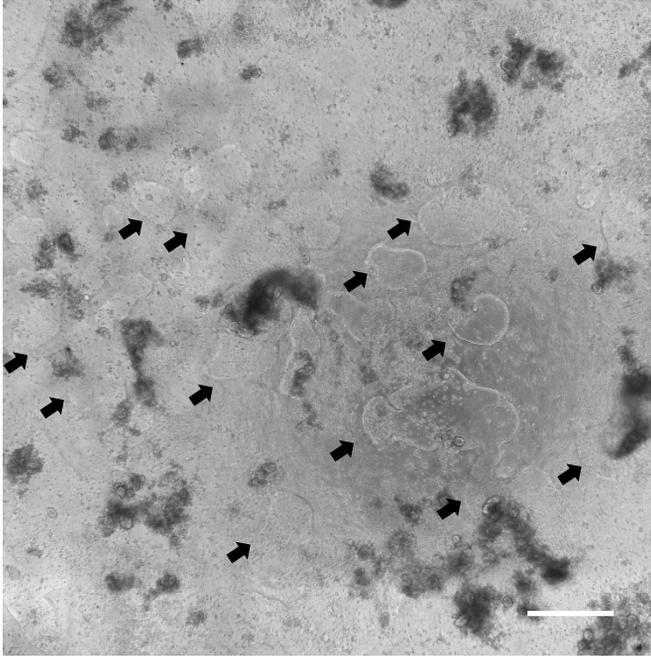
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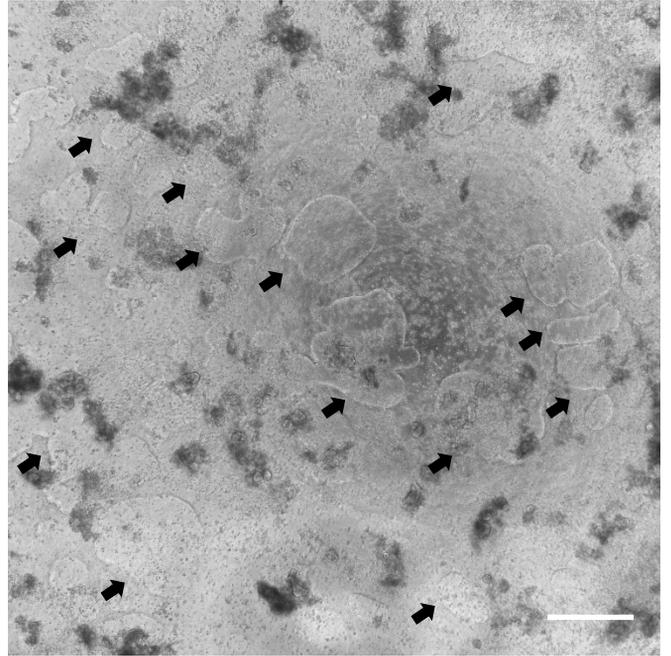
Supplemental Figure S1.

Characterization of rat duodenal organoid of cystic type. (a) Whole mount immunostaining of cystic type for Sox9 (green) and F-actin (white), Ki-67 (red) and F-actin (white), merge with DAPI (blue), CK20 (green) and P-gp (red) with DAPI (blue), ZO-1 (red) and DAPI (blue), phospho-Ezrin (p-Ezrin, red) and DAPI (blue). Scale bar = 100 μ m (b) Day 4 organoids sections were stained with HE, AB-PAS. Scale bar = 100 μ m. (c) Transmission electron microscopy of organoids on day 4. Arrows, desmosomes; MG, Matrigel; Mv, microvilli; Lu, Lumen. Scale bar = 2.0 μ m.

Well 1



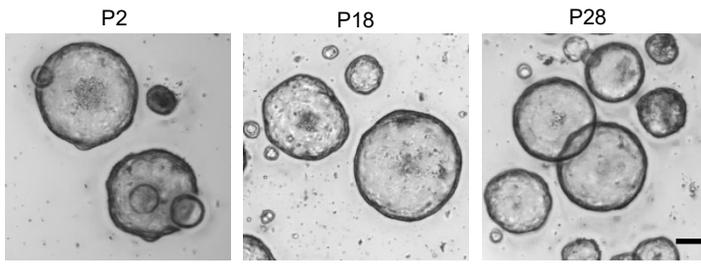
Well 2



Supplemental Figure S2.

Day 7 monolayer culture generated by conventional rat organoids. Arrows show the cell absence area.

Scale bar = 500 μm



Supplemental Figure S3.

Representative images of long-term cultured cyst-enriched organoids.

Scale bar = 100 μ m

Supplemental Table S1. Primers used for quantitative RT-PCR analysis

Gene	Forward primer	Reverse primer
<i>Si</i>	TGTTCCCTAAGGCTCGATGGTTTG	TTAGATGCGTGTTCCGTGCTG
<i>Abcb1a</i>	AGATCGTGAAGGCAGCCAAG	AGAAGTAAGATGTGAGGCTGTCTGA
<i>Abcb1b</i>	TGACTCGGGAGCAGAAGTTTGA	GGCGAAGGTGATCCCAAAGA
<i>Abcc2</i>	TGAGCACCAGCAGCGATTC	GAAGACGACCAAGTTTCCAACCA
<i>Cyp1a1</i>	GTCCCGGATGTGGCCCTTCTCAAA	TAACTCTTCCCTGGATGCCTTCAA
<i>Cyp3a9</i>	TTGTACGATGGTCGACAGCC	CCACTGGACCAAAGTTCCGT
<i>Act-b</i>	GGAGATTACTGCCCTGGCTCCTA	GACTCATCGTACTCCTGCTTGCTG