Modeling colorectal cancer using CRISPR-Cas9–mediated engineering of human intestinal organoids

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Human colorectal tumors bear recurrent mutations in genes encoding proteins operative in the WNT, MAPK, TGF-β, TP53 and PI3K pathways1–2. Although these pathways influence intestinal stem cell niche signaling3–5, the extent to which mutations in these pathways contribute to human colorectal carcinogenesis remains unclear. Here we use the CRISPR-Cas9 genome-editing system6,7 to introduce multiple such mutations into organoids derived from normal human intestinal epithelium. By modulating the culture conditions to mimic that of the intestinal niche, we selected isogenic organoids harboring mutations in the tumor suppressor genes APC, SMAD4 and TP53, and in the oncogenes KRAS and/or PIK3CA. Organoids engineered to express all five mutations grew independently of niche factors in vitro, and they formed tumors after implantation under the kidney subcapsule in mice. Although they formed micrometastases containing dormant tumor-initiating cells after injection into the spleen of mice, they failed to colonize in the liver. In contrast, engineered organoids derived from chromosome-instable human adenomas formed macrometastatic colonies. These results suggest that ‘driver’ pathway mutations enable stem cell maintenance in the hostile tumor microenvironment, but that additional molecular lesions are required for invasive behavior.

Multiple mutations or ‘hits’ are thought to be required for the transformation of normal colon epithelium into invasive and metastatic colorectal cancer (CRC). The adenoma–carcinoma sequence model proposes a molecular basis for carcinogenesis8: loss of the APC gene (encoding adenomatous polyposis coli) first induces a premalignant precursor lesion (an adenoma), and subsequent mutations in genes including KRAS (encoding Kirsten rat sarcoma viral oncogene homolog), SMAD4 (encoding SMAD family member 4) and TP53 (encoding tumor protein p53) promote transformation to an invasive and metastatic phenotype9. Mutations in these genes are thought to drive colorectal carcinogenesis by deregulating driver pathways that modulate cellular proliferation, differentiation and apoptosis in order to confer selective growth advantages to mutated epithelium, including the WNT, RAS-MAPK, PI3K, TGF-β and TP53 pathways6,7,10.

In rodent models, introducing combinations of two of these driver pathway mutations, such as Apc and Kras, Braf transforming gene (Braf) and Trp53 or Apc and Smad4, induces intestinal adenocarcinoma11–14. In contrast, each human CRC often harbors 3–6 recurrent driver mutations15, suggesting that higher numbers of driver mutations may be required for human colorectal carcinogenesis. Notably, the number of driver pathway alterations in human CRC is highly variable, with some CRCs carrying no pathway alterations or only one pathway alteration1–2. In these CRCs, somatic chromosomal aberrations or epigenetic lesions might instead alter driver pathway signaling. It is also possible that driver pathway alterations are dispensable in a subset of CRCs. To elucidate the role of driver pathway mutations in human CRC, a prospective analysis of the effects of site-directed mutations in the normal human colon epithelium is warranted, but this has long been hampered by the lack of a tractable culture system.

Recently, we developed an epithelial organoid culture system in which human intestinal stem cells (ISCs) indefinitely self-renew and form crypt-like organoid structures in Matrigel15,16. In this organoid culture system, human ISCs require a unique combination of Wnt, R-spondin, epidermal growth factor (EGF) and bone morphogenetic protein (BMP)/transforming growth factor (TGF)-ß inhibitors, which mimic conditions in the ISC niche. In a physiological setting, these factors are locally provided by neighboring epithelial and mesenchymal cells1,7,17, which restrict the self-renewal of ISCs to the narrow niche space at the crypt bottoms. In contrast, these niche factors are often dispensable for the growth of organoids derived from human CRC (as opposed to normal human epithelium)15. Because these niche factors influence most if not all of the pathways that are mutated in CRC, driver pathway mutations may modulate carcinogenesis by conferring selective growth advantages to ISCs18. As described here, to examine the influence of various driver pathway mutations in human colorectal carcinogenesis, we prospectively introduced multiple driver pathway mutations into human normal or adenoma-derived intestinal organoids.

To introduce defined genetic mutations in human intestinal epithelial cells, we used clustered regularly interspersed short palindromic repeat (CRISPR)–CRISPR-associated protein 9 (Cas9) genome editing6,7,19 in human intestinal organoids (Supplementary Fig. 1b).
and Supplementary Table 1). First, we generated CRISPR-Cas9 sgRNAs targeting the APC, SMAD4 and TP53 tumor suppressor genes (Fig. 1). Next, we dissociated human normal colonic organoids into single cells and electroporated the APC-targeting vectors into these cells; we then selected the cells in medium lacking Wnt and R-spondin. Under these culture conditions, single cells treated with control vector invariably ceased proliferation within 2–3 d, but the APC-targeted (A-) ISCs were able to form organoids (Fig. 1b and Supplementary Fig. 2a) in which mutations were detected by SURVEYOR (Transgenic) assay. After clonal expansion of the single-cell derived A-organoids, the disruption of the targeted locus was confirmed by sequence analysis (Fig. 1a). As expected, western blotting showed the upregulation of β-catenin (Supplementary Fig. 2b). Next, we used similar methods to target SMAD4 in dissociated normal organoids. As previously reported, normal intestinal organoids require noggin and a TGF-β receptor inhibitor (A83-01) for sustained growth. Thus, we selected SMAD4-targeted (S-) organoids in a medium lacking noggin and A83-01 but containing the MDM inhibitor nutlin-3. (Supplementary Fig. 2c) and KRAS-targeted (T-) but not control organoids were readily recovered from cultures in medium containing nutlin-3, an inhibitor of MDM2 (ref. 21) (which degrades TP53) (Fig. 1f). Cloned T-organoids consistently showed indel mutations in the targeted locus (Fig. 1e and Supplementary Fig. 2c) and overexpressed a truncated stabilized form of TP53 (Supplementary Fig. 2f).

By using the genome-editing system established above, we next targeted KRAS that is downstream of EGF signaling by introducing the KRAS<sup>G12V</sup> hot spot mutation into A-organoids. To knock in the KRAS<sup>G12V</sup> mutation, we electroporated a sgRNA targeting a sequence adjacent to KRAS exon 2 together with a donor vector containing homologous arms but encoding the KRAS<sup>G12V</sup> mutation (Supplementary Fig. 2d). Removal of EGF or addition of an EGF receptor (EGFR) inhibitor to

Figure 1 CRISPR-Cas9–mediated introduction of driver pathway mutations in human intestinal organoids. (a–j) Human intestinal organoids were dissociated into single cells and electroporated with sgRNAs targeting APC, SMAD4 or TP53. (a,c,e) Sequences of targeted loci. Note that the splicing donor site is disrupted in one allele of APC. (b) Untargeted wild type (WT) or APC-targeted (A-) organoids were established and dissociated into single cells. The single cells were cultured in control medium (containing Wnt, EGF, noggin, R-spondin and A83-01) or medium lacking Wnt and R-spondin but containing EGF, noggin and A83-01. (d) Control or SMAD4-targeted (S-) organoids were cultured in control medium or medium lacking noggin and A83-01 but containing TGF-β. (f) Control or TP53-targeted (T-) organoids were cultured in control medium or medium containing the MDM inhibitor nutlin-3. (g) Strategy for targeting the human KRAS locus. The donor vector harbors the KpnI site (light blue) and a G12V mutation (red) was co-electroporated to A-organoids. Sequences of six mutant alleles (Del) from targeted organoid clones are shown. Ref, reference sequence. Two of these clones show partial knock-in with wild-type codon 12 (WT). In the other four alleles, the homologous region is completely knocked in (KI). E: EcoRI, K: KpnI. (h) Control KRAS<sup>WT</sup> A-organoids or KRAS<sup>G12V</sup> A-organoids were grown in control medium or medium lacking EGF. (i) Strategy for targeting the human PIK3CA locus. The donor ssDNA contains the E545K mutation (red). (j) KRAS<sup>G12V</sup> or KRAS<sup>G12V</sup>PIK3CA<sup>E545K</sup> A-organoids were cultured in medium lacking EGF, or medium lacking EGF and containing a MEK inhibitor (MEKi). The results are representative of three different experiments performed in triplicate (b,d,f,h,j). Scale bars, 1 mm (b,d,f,h,j).
the culture medium blocked colony formation by untargeted ISCs, enabling the enrichment of organoids derived from KRAS<sup>G12V</sup>-targeted ISCs (Fig. 1h). Homologous recombination in cloned KRAS<sup>G12V</sup> organoids was confirmed by Southern blotting and sequence analysis (Fig. 1g and Supplementary Fig. 2g). As expected, the efficiency of the mutation knock-in was lower than that observed after the introduction of indel mutations (Supplementary Table 2). Constitutive activation of KRAS was verified by ERK phosphorylation in KRAS<sup>G12V</sup> organoids in the absence of EGF (Supplementary Fig. 2h). Notably, the KRAS<sup>G12V</sup> mutation was not sufficient to fully substitute for EGFR signaling, as the removal of EGF decreased the growth efficiency of the KRAS<sup>G12V</sup> organoids (Fig. 1h). We next used CRISPR-Cas9 to activate another downstream signaling component of EGF—the PI3K-AKT pathway—in the KRAS<sup>G12V</sup> organoids. A sgRNA targeting a locus adjacent to the mutational hot spot (PIK3CA<sup>E545K</sup>) was electroporated into dissociated KRAS<sup>G12V</sup> organoids with a homologous donor oligonucleotide carrying the mutation (Fig. 1i). The PIK3CA<sup>E545K</sup> knock-in (KRAS<sup>G12V</sup>PIK3CA<sup>E545K</sup>) organoids selectively grew in the absence of EGF, in the same selection culture conditions used for KRAS<sup>G12V</sup> organoids (Fig. 1j). Addition of a mitogen/extracellular signal-regulated kinase (MEK) inhibitor, which blocks the downstream signaling of KRAS, to the culture medium—enriched KRAS<sup>G12V</sup>PIK3CA<sup>E545K</sup> organoids from KRAS<sup>G12V</sup> (Fig. 1j). Recovered cloned organoids showed correct homologous recombination, which was confirmed by sequence analysis (Fig. 1i), as well as by phosphorylation of AKT (Supplementary Fig. 2i), an indicator of constitutive activation of PI3CA. In contrast to the single KRAS<sup>G12V</sup> mutation, the KRAS<sup>G12V</sup>PIK3CA<sup>E545K</sup> double mutations completely substituted for EGF signaling (Fig. 1j).

By using this genome editing and selection system for single-cells derived from organoids, we generated isogenic organoids carrying various combinations of driver gene mutations recapitulating the adenoma-carcinoma sequence (Fig. 2a and Supplementary Fig. 1c). We found that each culture condition was strictly specific to the corresponding pathway mutation, which enabled us to introduce various combinations of the pathway mutations in organoids without using conventional antibiotics-based selection (Supplementary Fig. 1a). After multiple cycles of cloning and CRISPR-Cas9-mediated targeting, engineered organoids progressively decreased the niche factor requirements (Fig. 2b). Accordingly, A-organoids were sensitive to treatment with EGFR, TGF-β, nutlin-3 or MEK inhibitors. With this antibiotic-free approach, we established AKSTP-organoids carrying

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**Figure 2 Sequential engineering of various combinations of driver mutations.** (a) Strategy for establishing multiple driver mutations—to mimic the adenoma-carcinoma sequence—in organoids using CRISPR-Cas9. Engineered organoids were selected by the indicated niche factor–modulated culture conditions. W, Wnt-3A; R, R-spondin-1; E, EGF; N, noggin; A, AB3-01; Nut, nutlin-3; TGF, TGF-β, MEK inhibitor. Detected mutations (A, APC; K, KRAS<sup>G12V</sup>; T, TP53; S, SMAD4; P, PI3KCA<sup>E545K</sup>) are shown. Sequences are shown in Supplementary Figure 8a. (b) Driver gene mutations provide resistance to niche-modulated culture conditions. The results are representative of two different experiments performed in triplicate. (c) Immunostaining of engineered organoids. Left, F-actin stained by phalloidin (green) integrin-α6 (red). Right, MUC2, mucin 2 (green); ChgA, chromogranin A (red). Nuclear counterstain: Hoechst 33342 (blue). Scale bars, 50 μm. Results are representative of two different experiments. (d) Copy number analysis in engineered organoids. Blue and red indicate gain and loss of chromosome, respectively. See Supplementary Figure 4 for detail of BKSTP. (e) Gene set enrichment analysis using CRC data sets (90 genes). Enrichment score = 1.721, False discovery rate = 0.003. (f) Principal component (PC) analysis of engineered organoids (blue), adenoma organoids (red) and CRC organoids (brown). (g) Heat maps showing the average log<sub>2</sub>-fold difference of the indicated genes between the engineered organoids and CRC organoids. See Supplementary Figure 3 for heat maps including adenoma organoids.

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five gene mutations (APC, KRAS (G12V), SMAD4, TP53 and PIK3CA (E545K)), which grew completely independently of all niche factors described above (Fig. 2b). The engineered organoids were largely devoid of aneuploidy, copy number alterations (Fig. 2d), or CpG-island methylator phenotype (CIMP) (not shown), indicative of the maintenance of genetic stability during the course of organoid culture and gene manipulation. In addition to their nearly normal genetic status, whole-mount immunohistochemistry revealed intact apico-basal polarity and multiple features of well-differentiated colonic epithelium within the engineered organoids (Fig. 2c).

We next investigated the effect of driver pathway mutations in human intestinal epithelium on global gene expression patterns. Five lines of adenoma organoids and nine lines of CRC organoids were established and used for microarray analysis (Supplementary Table 1).

To determine whether gene signatures of organoids represented their original tumor stage, we used adenoma and CRC gene expression data sets obtained from patients\(^2\). Gene set enrichment analysis (GSEA) indicated that the gene expression alteration during tumor progression was largely retained after organoid culture despite differences in micro-environments (Fig. 2c). Principal component analysis (PCA) among the three groups of organoids (CRC, adenoma and engineered) revealed similarity between the engineered and adenoma organoids (Fig. 2f). In the genes differentially expressed between engineered and CRC organoids, the engineered organoids were enriched for genes related to secretory differentiation (for example, MUC2, SPINK4, TFF1 and CHGA), which are often lost in the course of colorectal carcinogenesis but retained in adenoma organoids\(^2\) (Fig. 2g and Supplementary Fig. 3).

By contrast, CRC-specific genes (for example, EREG, TRPM2 and

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**Figure 3** Engraftment of engineered, adenoma and CRC organoids in NOG mice. Indicated EGFP-labeled organoids were implanted under the kidney capsule of NOG mice. (a) Photographs of four representative tumors isolated 1 month (1M) or 2 months (2M) after the transplantation. (b) Average surface area of xenografts derived from organoids at 1 or 2 months after transplantation (n = 4 at each time point). The value of CRC is the average of six individual lines of CRC organoids (n = 24). Error bars indicate sem, * P < 0.05. (c) In situ hybridization of LGR5 mRNA (brown) in xenografts derived from organoids. (d) Representative HE staining of kidney subcapsule xenografts derived from AKSTP-organoids isolated 1 month after transplantation. Insets are magnified at bottom. The inset in the right panel indicates region shown in f. (e) Left, immunostaining of α-smooth muscle actin (SMA) in xenografts derived from AKSTP-organoids. Right, co-immunostaining of EGFP (green, human-derived organoids) and SMA (red). Nuclear counterstaining with Hoechst 33342 is shown in blue. (f) Immunohistochemistry of AKSTP tumors. β-catenin; ChgA, chromogranin A; KRT20, cytokeratin 20. (g) Representative pictures of tumors that metastasized to the liver after injection into the spleen of NOG mice (n = 4 mice per genotype). Livers with metastatic lesions 2 months after the injection are shown. Top images show bright-field images of the hepatic lobule. Insets show bright-field images of the whole liver. Bottom images show an EGFP-fluorescent metastasized tumor in the hepatic lobule. Arrowheads indicate micrometastatic lesions. (h) H&E and immunostaining of metastasized organoids in the liver. (i) Tumor formation of organoids derived from AKSTP-micrometastatic lesions (1 month after transplantation). (j) Representative (of three images) H&E staining of tissue shown in i. Scale bars, 5 mm (a,g,i), 1 mm (d, top), 100 μm (h), 50 μm (c,d (bottom), e,f,h,j).
ETV4) were not expressed in the engineered organoids. Although the biological impact of these gene expression changes remains unknown, our results suggest that driver pathway mutations and subsequent transcriptional changes are insufficient for the gene expression re-programming that occurs during the transition to CRC.

To determine the role of driver mutations in human colonic tumorigenesis, we transplanted GFP-labeled organoids into kidney subcapsules of Nod−/−IL2ręnull (NOG) mice (Fig. 3a). The CRC organoids efficiently formed tumors that mimicked their parental histology, but the A-organoids or adenoma organoids showed no or minimal engraftment, suggesting that the kidney subcapsule is an unfavorable niche for benign epithelium (Fig. 3a,b). The engineered organoids acquired tumorigenic capacity in proportion to the number of mutations they harbored, as AKST- or AKSTP-organoids formed readily visible tumors (Fig. 3a,b). Isolated AKST tumors were re-propagated in vitro without niche factors (not shown). The tumor-derived AKST-organoids retained their original engineered mutations and could be re-engrafted under the kidney subcapsules of NOG mice (not shown). Consistent with their tumorigenic capacity, leucine-rich repeat–containing G protein–coupled receptor 5 (LGR5) stem cells were detected in growing tumors from AKST and AKSTP-organoids but not in remnants of engrafted adenoma organoids (Fig. 3c). We observed a similar tumorigenic capacity of engineered organoids derived from a second independent individual (AKST′ in Fig. 3b and Supplementary Fig. 1).

To assess off-target cleavage by CRISPR-Cas9, we examined potential off-target sites in AKSTP-organoids by exome sequencing (Supplementary Table 3). By using computer-based off-target prediction, we found that there were nine off-target sites carrying three base pair mismatches, of which six contained no mutations and the others were in the 3′UTR or intronic regions (Supplementary Table 3). We also analyzed 94 off-target sites with four mismatches, of which 70 were covered by exome sequencing without detecting mutations, and the others were in non-exonic regions (Supplementary Table 3). To further validate that the phenotypes we observed were due to specific mutations introduced by CRISPR-Cas9, we engineered organoids using lentiviruses encoding shRNA to knock down SMAD4 and TP53 as well as transgenes encoding CTNNB1 (constitutively active form of β-catenin to activate WNT signaling), KRASG12V and PIK3CAE545K (Supplementary Fig. 4). The resulting organoids, termed BKSTP-organoids, exhibited identical in vitro and in vivo phenotypes as those of AKSTP-organoids (Supplementary Fig. 5). These results suggest that driver gene mutations cooperatively substitute for niche signaling and confer selective growth advantages to SCs in unfavorable microenvironments.

Histological analyses of AKST/AKSTP and BKSTP tumors revealed adenoma histology represented as well-ordered, highly differentiated epithelium encapsulated by mouse-derived stromal cells (Fig. 3d–f and Supplementary Figs. 5 and 6). Although AKSTP tumors had partial features of low-grade adenocarcinoma (Fig. 3d), their non-invasive
phenotype restricted their growth after 2 months. By contrast, most of the CRC organoids invaded adjacent tissue, resulting in constant enlargement of the tumor mass over 2 months (Fig. 3b and Supplementary Fig. 6). The functional difference between engineered and CRC organoids was even more evident when considering their metastatic ability. When injected into the mouse spleen, organoids derived from metastatic CRC successfully metastasized and colonized the liver, whereas AKSTP-organoids only formed micrometastatic lesions (<1 mm²) (Fig. 3g and Fig. 4e). Histological analysis of AKSTP micrometastases depicted differentiated epithelium largely devoid of proliferation (Fig. 3h). Notably, cells from AKSTP micrometastases were capable of forming tumors after retransplantation into the kidney subcapsule, indicating that the micrometastatic lesions harbored dormant tumor-initiating cells (Fig. 3i, j). These results suggest that combinations of recurrent driver gene mutations contribute to niche-independent stem cell maintenance but not to metastatic progression.

In the adenoma-carcinoma sequence model, cytogenetic lesions are thought to drive tumor progression. However, control of chromosomal rearrangement of human primary cells is technically unfeasible. We engineered three lines of adenoma organoids (Fig. 4a and Supplementary Figs. 1, 7–9) one of which showed a chromosomal instability (CIN) phenotype during the CRISPR-based engineering of TP53 and SMAD4 (hereafter referred to as AdɛCIN TS-organoids) (Fig. 4a, b). Similarly to parental adenoma organoids and AKSTP-organoids, AdɛCIN TS-organoids displayed highly polarized morphology and failed to macro-metastasize to the liver after injection into the spleen (Fig. 4c–e). However, after we used CRISPR-Cas9 to introduce a KRASG12V mutation (AdɛCIN TSK-organoids) or lentivirus to introduce a PIK3CAE545K mutation (AdɛCIN TSP-organoids), these further-engineered AdɛCIN TSK/TSP-organoids showed cribriform-like dysplastic structures in vitro (Fig. 4c) and were capable of forming large metastatic tumors displaying histologic malignancy (Fig. 4d–f). A microarray analysis revealed no global gene expression changes during the metastatic transformation of the adenoma organoids (Supplementary Fig. 7). We confirmed no predicted off-target mutations in AdɛCIN TSK-organoids (Supplementary Table 3). Of note, CIN-negative adenoma remained non-metastatic after CRISPR-TSK or lentivirus-TSK engineering (Supplementary Fig. 9).

Although the molecular mechanism of the metastatic progression remains unknown, these results suggest that genetic lesions other than driver pathway mutations are required for colon carcinogenesis.

The CRISPR-Cas9 system has been used to generate cancer cells from mouse primary cells or directly from in vivo tissue. Here we established an efficient system to introduce multiple driver gene mutations into the human colonic epithelium by niche factor-modulated culture conditions, enabling clonal expansion of mutant stem cells without antibiotic selection. By using this niche-based selection system, we obtained quintuple pathway-mutant organoids, which grew independently of niche factors and formed tumors in hostile niche environments. Notably, most of CRC (>90%) as well as genetically engineered mouse CRC models possess fewer than five driver pathway alterations, suggesting that the non-mutated niche signaling in CRC could be complemented by adjacent non-epithelial mesenchymal or immune cells. Although driver pathway mutations increased the robustness of stem cells by hijacking niche signaling, driver pathway mutations per se were insufficient for malignant progression in this system. Interestingly, our results implicate chromosomal instability, in addition to driver pathway mutations, as required for invasive and metastatic transformation. Further studies will be required to elucidate the functional contribution of other molecular lesions, such as aneuploidy, copy number variations or epigenetic alterations in the malignant progression of human CRC in this system.

METHODS

Methods and any associated references are available in the online version of the paper.

Accession codes. Gene expression data have been deposited in the NCBI Gene Expression Omnibus with accession number GSE57965.

Note: Any Supplementary Information and Source Data files are available in the online version of the paper.

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AUTHOR CONTRIBUTIONS

M.M., S.D., M.S., A.T., M.F. and Y.O. performed experiments. T.W. and T.K. provided biological specimens. T.S. conceived and designed the project. T.S. wrote the manuscript.

COMPETING FINANCIAL INTERESTS

The authors declare competing financial interests: details are available in the online version of the paper.

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ONLINE METHODS

Culture of human intestinal organoids. Surgically resected intestinal tissues or endoscopic biopsy samples were obtained from Tokyo University Hospital and Keio University Hospital. This study was approved by the Keio University Ethics Committee and Human Genome, Gene Analysis Research Ethics Committee, Graduate School of Medicine, the University of Tokyo. All samples were obtained from patients who provided informed consent before surgery or endoscopy. Adenoma, CRC and normal epithelia were isolated and cultured according to previously reported protocols. The basal culture medium for human intestinal organoids was slightly modified from our original report as follows: Advanced Dulbecco’s modified Eagle medium/F12 was supplemented with penicillin/streptomycin, 10 mM HEPES, 2 mM GlutaMAX, 1 x B27 (Life Technologies), 10 mM gastrin I (Sigma) and 1 mM N-acetylcysteine (Sigma). The following niche factors were used: 50 ng/ml mouse recombinant EGF, 100 ng/ml mouse recombinant noggin (Peprotech), 10% R-spondin-1 conditioned medium (kindly provided by C. Kuo, Stanford University)34, 50% Wnt-3A conditioned medium35, 500 nM A83-01 (Tocris) and 10 µM SB202190 (Sigma). To select mutant organoids, the following reagents were used: 10 ng/ml mouse recombinant TGF-β (R&D), 100 ng/ml mouse recombinant BMP4 (Peprotech), 1 µM EGFR inhibitor (EMD Millipore), 50 nM PD325901 (MEK inhibitor) (Sigma) or (+)-nuttlin-3 (Cayman Chemical). For the organoid formation assay and before genome editing, organoids were dissociated into single cells with TrypLE Express (Life Technologies) and 1,000 cells were cultured in a 48-well plate under the above culture conditions for 10 d. To prevent anoxia, 10 µM Y-27632 was included in the culture medium for the first 2 d (ref. 3). The growth of organoids was estimated by area of intact organoids using bioimaging software (LuminaVision, Mitani Visual).

Genome engineering. Gene-specific sgRNA oligos were cloned into the px330 expression vector29 (Addgene), which bicistronically expresses sgRNA and Cas9 nuclease. The sgRNA sequence was determined by the CRISPR Design Tool (http://crispr.mit.edu/) (Supplementary Table 4). After dissociating the organoids with TrypLE express, cells were electroporated with 10 µg of CRISPR-Cas9 plasmid (NEPpgane). Cells were washed, embedded in Matrigel and cultured with optimal growth factors and Y-27632 (Sigma). For xenotransplantation assay, 7 µg of the piggyBac GFP-Puro expression vector and 3 µg of the PBase vector (SBI) were electroporated to fluorescently label the organoids. To knock-in an oncogenic mutation, 5 µg of sense or antisense donor soLoigo (manufactured by Life Technologies) or 5 µg of homologous recombination donor plasmid (constructed from pCR2.1 TOPO vector) was co-delivered with the sgRNA/Cas9 plasmid. Genomic DNA extracted from engineered organoids was used for the SURVEYOR assay (Mutation Detector kit, Transgenomics)35 or Southern blotting analyses. For validation of targeted mutations, genomic DNA was isolated from cloned engineered organoids, followed by PCR amplification of targeted loci. PCR products were directly sequenced or cloned into a TOPO-TA cloning vector according to the manufacturer’s instructions (Life Technologies). Plasmid DNA was isolated from eight bacterial clones and grown biallelic gene alterations. For whole-exome sequencing, genomic DNA was isolated from cloned engineered organoids, followed by PCR amplification using bioimaging software (LuminaVision, Mitani Visual).

Western blot analysis. The Matrigel surrounding the organoids was depolymerized by a Cell Recovery Solution (BD Biosciences). The released organoids were washed and pelleted. The pelleted organoids were lysed with Cell Lysis Buffer (CST) containing a protease/phosphatase inhibitor cocktail (Thermo Science) and phosphatase inhibitor cocktail set V (Calbiochem), and the organoids were further disrupted by sonication. Equal amount of SDS sample buffer was applied to the cell lysates. Protein was quantitated by the Bradford assay, and equal amounts of protein were loaded in each lane of a 4–20% SDS–PAGE. Standard western blot analyses were performed using the following primary antibodies: anti-β-catenin (610153, BD Biosciences, 1:1,000), anti-β-actin (A1978, Sigma, 1:1,000), anti-SMAD4 (sc-7966, Santa Cruz, 1:1,000), anti-TP53 (sc-98, Santa Cruz, 1:1,000), anti-ERK-1/2 (#9102, CST, 1:1,000), anti-phospho T202/Y204 ERK-1/2 (#4370, CST, 1:1,000), anti-AKT-1/2/3 (#4072, CST, 1:1,000), anti-phospho S473AKT (#9271, CST, 1:1,000). Signals were detected by HRP-conjugated secondary anti-rabbit antibody and visualized with the ECLSelect LASS500 (GE Healthcare) and Image Quant LAS 500 (GE Healthcare).

Microarray and copy number analysis. Tumor organoids were expanded for gene expression analysis or copy number analysis. For gene expression microarray analysis, 300 ng of whole RNA was extracted from organoids using the RNeasy Mini Plus kit (Qiagen). Microarray hybridization was performed on the GeneChip Prime View Human Gene Expression Array (Affymetrix) according to the manufacturer’s instructions. The data were analyzed with GeneSpring GX (Agilent). The microarray data are available in the GEO database (Accession number: GSE57965). GSEA was performed using open source software (v.2.0, Broad Institute, http://www.broadinstitute.org/gsea/index.jsp). A ranked gene list was created between adenoma and CRC organoids, and it was computed with a published pre-ranked list of upregulated genes in patient CRC specimens compared to those of adenoma.22 The number of permutations was 1,000. For copy number analysis, the genomic DNA of organoids was extracted using the QIAamp DNA Blood Mini Kit (Qiagen) and analyzed by CytoScan HD (Affymetrix). The copy number data were analyzed by Nexus copy number software (BioDiscovery).

Xenotransplantation of organoids. NOD.Cg-Prka cm1SugIl2rg tm1Sug/Ctr j (NOG) mice (male, 7–12 weeks old) were obtained from the Central Institute for Experimental Animals (CIAE, Japan) and housed under specific pathogen–free conditions. All procedures were approved by the Keio University School of Medicine Animal Care Committee. The GFP-expressing organoids were isolated from Matrigel using Cell Recovery Solution (BD Biosciences) and mechanically dissociated into small clusters of cells. The cell clusters (equivalent to 1 x 10^5 cells) were resuspended in cold Matrigel and injected under the kidney subcapsules or spleens of NOG mice. One or two months after transplantation, the kidney or liver was isolated, and the tumor size was measured by area of GFP fluorescence (Nikon Multi-zoom microscope, LuminaVision software). A blinded investigator performed the measurement of the tumor size. The grafts were fixed for subsequent histological analysis. In some xenografts, tumor cells were isolated and subsequently cultured according to the methods described above. Each line of engineered organoids or CRC organoids was transplanted into 8 independent NOG mice. Four mice were used for analysis at each time point. The sample size was determined as the minimum size necessary to obtain a significant difference (P = 0.05) at a power of 80% when there was a 30% change in the tumor size. The time point of euthanasia was randomly assigned. As for inclusion/exclusion criteria for the xenograft experiments, we pre-determined to exclude data from mice euthanized because of sickness. We euthanized 2 out of 60 mice with implanted CRC organoids due to the tumor burden during the experimental period (2 months) and excluded these mice from the data analysis.

Immunohistochemistry and in situ hybridization. Whole-mount immunostaining was performed as previously described. Isolated xenografts were immediately fixed with 4% paraformaldehyde. The standard protocol for sectioning paraffin-embedded tissues and H&E or Periodic acid–Schiff (PAS) staining was used. For immunohistochemistry, the following primary antibodies were used: anti-integrin-α6 (313601, Biolegend, 1:100), anti-Ki67 (RM-9106-S, Thermo Scientific, 1:300), anti-cytokeratin-20 (NCL-L-CK20, Leica, 1:50), anti-α-smooth muscle actin (MS-113-P, Thermo Scientific, 1:800), anti-EGFP (Ab6673, Abcam, 1:500), anti-Mucin2 (sc-15334, Santa Cruz, 1:100), anti-chromogranin A (sc-1488, Santa Cruz, 1:100) and anti-β-catenin (610153, BD Biosciences, 1:100). Secondary antibodies were polymer HRP conjugated anti-rabbit or mouse antibody (Nichirei Bioscience) for paraffin sections; Alexa Fluor 488 or 568 conjugated anti-rabbit, rat or goat antibody (Life Technologies) for whole-mount staining. Alexa Fluor 647–phalloidin (A22287, Life Technologies) was used for the visualization of F-actin. Nuclei were counterstained with hematoxylin (paraffin sections) or Hoescht 33342 (whole mount). For paraffin sections, staining was developed with DAB+ substrate chromogen (Dako). In situ hybridization for LGR5 mRNA was performed with RNA Scope (Advanced Cell Diagnostics).
according to the manufacturer's instructions. The images of organoids were taken by either confocal microscopy (Leica SP8) or digital microscope (Keyence).

**Lentiviral transduction.** The VSV-G–pseudotyped third generation self-inactivating lentiviral vector was used for gene knockdown or over-expression. CTNNB1<sup>S33Y</sup>, KRAS<sup>G12V</sup> and PIK3CA<sup>E545K</sup> were cloned into the pLenti CMV GFP Neo/Hygro/Puro/Zeo plasmid (Addgene). miR sequences targeting SMAD4 and TP53 were designed using The CRISPR Design Tool ([http://crispr.mit.edu/](http://crispr.mit.edu/)) and inserted into the pDNA6.2-GW/EmGFP-miR vector (BLOCK-iT Pol II mi RNAi kits, Life Technologies) ([Supplementary Table 4](#)). The miR vectors reduced the target gene expression by at least 80% in HEK293 cells. The viral vectors were transfected into HEK293FT cells with third-generation packaging plasmids (pLP1, pLP2 and pLP/VSVG; Invitrogen). The viral supernatant was passed through a 45-µm filter and centrifuged at 6,000 g overnight. The titer of the viral supernatant was determined by qPCR and was typically higher than 10<sup>7</sup> titer units per ml. The organoids were dissociated into single cells and transduced with lentivirus as previously described.<sup>29</sup> Three days after the transduction, the infected organoids were selected with the appropriate antibiotics as follows: G418 (100 µg/ml), hygromycin (15 µg/ml), puromycin (2 µg/ml) and zeocin (200 µg/ml).

**Statistical analysis.** All results are expressed as mean ± sem and contain a minimum of n = 4 per group. We used unpaired Student’s t-test or the analysis of the xenograft experiments. For gene expression microarray analysis, we used the Storey bootstrapping method. P < 0.05 was considered statistically significant. The experiments were not randomized.

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