

AUS DEM DEUTSCHEN KREBSFORSCHUNGSZENTRUM
UND DEM LEHRSTUHL FÜR ZELL- UND MOLEKULARBIOLOGIE
DER MEDIZINISCHEN FAKULTÄT MANNHEIM
LEITUNG: PROF. DR. MICHAEL BOUTROS

Multiparametric phenotyping of intestinal organoids to model disease initiation and treatment response in colorectal cancer

INAUGURALDISSERTATION

ZUR ERLANGUNG DES MEDIZINISCHEN DOKTORGRADES
AN DER
MEDIZINISCHEN FAKULTÄT HEIDELBERG DER
RUPRECHT-KARLS-UNIVERSITÄT

VORGELEGT VON NIKLAS TIMON RINDTORFF
AUS TÜBINGEN

2022

DEKAN: PROF. DR. ANDREAS DRAGUHN
DOKTORVATER: PROF. DR. MICHAEL BOUTROS

I DEDICATE THIS THESIS TO ANDREA AND KLAUS RINDTORFF AS WELL AS SOPHIE WEIL. YOU
SUPPORTED ME THROUGH THE DARKEST AND BRIGHTEST HOURS OF THIS JOURNEY.

FIGURES WITHIN THIS THESIS WERE CREATED USING BIORENDER.COM

Contents

1	INTRODUCTION	15
1.1	Disclosure	15
1.2	The Colon	16
1.3	Colorectal Cancer	19
1.4	Colorectal Cancer Therapy	23
1.5	Cancer Drug Discovery	24
1.6	Multi-View Representation Learning in Biology	25
1.7	Towards causal Representation Learning in biology	25
1.8	Aims of Thesis	25
2	MATERIALS AND METHODS	29
2.1	Patients	29
2.2	Organoid Culture	30
2.3	Biochemical assays	32
2.4	Organoid profiling	35
2.5	Image analysis	38
3	IMAGE-BASED PROFILING OF ORGANOIDS	41
3.1	Disclosure	41
3.2	Establishing patient derived organoids for image-based profiling	42
3.3	Enabling methods for high-throughput image-based profiling of organoids	45
4	PROFILING OF ORGANOIDS IDENTIFIES MOLECULAR DETERMINANTS OF CANCER ORGANOID ARCHITECTURE AND PLASTICITY	49
4.1	Disclosure	49
4.2	Image-based profiling captures the morphological diversity of patient-derived cancer organoids	50
4.3	Organoid phenotype-profiles capture organoid viability	55
4.4	Drug induced organoid phenotypes correspond to drug mechanism of action	58
4.5	Multi-omics factor analysis identifies shared factors linking morphology, genomic data and drug activity	63
4.6	An IGF1R signaling program is associated with increased organoid size, decreased EGFR inhibitor activity and can be induced by mTOR inhibition	66
4.7	An LGR ₅ + stemness program is associated with cystic organoid architecture and can be induced by inhibition of MEK	70

5	PROFILING TO IDENTIFY REVERTANT THERAPEUTICS IN PRE-MALIGNANT MODELS OF COLON CANCER	74
5.1	Motivation	74
5.2	Generation of organoid colon adenoma models	76
5.3	Molecular profiling of organoid models	79
5.4	Image-based profiling of organoid models	83
5.5	Quantifying small molecule induced phenotypes across organoid models	85
5.6	Multi-omics factor analysis identifies shared factors linking functional and structural biological views	87
5.7	A canonical Wnt signaling associated program caused by Apc loss	88
5.8	An oncogene-induced senescence program caused by isolated KrasG12D activation	94
5.9	oncogenic Kras leads to increased mTOR signaling in the context of Apc loss of function	97
5.10	Ellagic acid is a candidate small molecule revertant therapeutic of organoid adenoma states	99
6	DISCUSSION	101
	APPENDIX A SOME EXTRA STUFF	iv
	REFERENCES	ix

Listing of figures

1.1	Colon Cancer Progression	20
1.2	21
1.3	27
3.1	Core liquid handling methods a Organoid isolation procedure. Colorectal cancer tissue biopsies were collected via endoscopy, enzymatically removed from extracellular matrix proteins, washed and resuspended in basal membrane extract hydrogel. After solidification of hydrogel domes, organoids were overlayed with growth factor rich culture medium. b Organoid high-throughput experimentation. Colorectal cancer organoids were harvested, partially digested, seeded in hydrogel-coated 384-well plates	42
3.2	Organoid cohort overview a Tumor location (right/left/rectum) and AJCC/UICC stage of colorectal cancers that patient derived organoids were derived from. b Consensus molecular subtypes of organoids determined by RNA expression analysis. c Mutation status in PDOs, as analyzed by amplicon sequencing. Figure created with support from Johannes Betge (graphical presentation), Erica Valentini (sequencing data analysis) and Benedikt Rauscher (CMS type inference). Figure adapted from <i>The drug-induced phenotypic landscape of colorectal cancer organoids</i> ³	44
3.3	Overview of experiments a Organoids were isolated from endoscopic biopsies from patients with colorectal cancer. Organoids were dissociated and evenly seeded in 384-well plates before perturbation with an experimental (464 compounds) and a clinical compound library (63 compounds à 5 concentrations, 842 perturbations across both libraries). After treatment, high-throughput fluorescence microscopy was used to capture the morphology of organoids. The multi-channel (DNA, beta-actin, cell permeability) 3D imaging data was projected, segmented, and descriptive features were extracted to quantify potential drug-induced phenotypes. Untreated organoid morphology, organoid size and drug activity scores were integrated with mRNA expression and mutation data in a Multi-Omics Factor Analysis (MOFA) to increase interpretability of organoid variation. Figure created with support from Johannes Betge (graphical presentation) and adapted from <i>The drug-induced phenotypic landscape of colorectal cancer organoids</i> ³	45

- 3.4 Image-based profiling method a The image-processing pipeline illustrated with representative example images from two organoid lines: Organoids were imaged at multiple layers along the z-axis. Images were projected using a maximum contrast projection and segmented using a convolutional neural network, both designed and implemented by Jan Sauer. Descriptive features were extracted from all three channels to quantify phenotypes. Feature plots show the median phenotype of unperturbed organoids, six example features (Area, Phalloidin intensity, DAPI intensity, FITC intensity, FITC Haralic angular second moment (ASM) and FITC intensity 1-percentile) and their z-scores relative to all profiled organoid lines are shown. Figure created with support from Jan Sauer (data processing) and Johannes Betge (graphical presentation). Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³ 46
- 4.1 Dataset dimensions and compound library overview a Number of organoid models and number of perturbations in previous publications reporting high-throughput drug screenings with patient derived cancer organoids, b Graphical representation of the compound libraries used for drug screening in this project: A library targeting kinases and stem cell pathways (KiStem library, 464 compounds) and a clinical library with 63 drugs in 5 concentrations. Figure created with support from Johannes Betge and adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³ 50
- 4.2 Image-based profiling captures the phenotype diversity of patient derived cancer organoids a Uniform Manifold Approximation and Projection (UMAP) of organoid-level features for a random 5% sample out of ca. 5.5 million organoids. The same sample is used for visualizations throughout the figure. Color corresponds to the log-scaled organoid area (dark blue: minimum size, yellow: maximum size). b organoid size distribution across lines. c UMAP representation of DMSO treated and drug treated organoids. Graph-based clustering of organoids by morphology. d UMAP embeddings of selected organoid lines (baseline state / 0.1% DMSO control-treated organoids) representing different morphological subsets, grey background consists of randomly sampled points. Depicted are representative example images for each line (right, cyan = DNA, magenta = Actin, scale-bar: 200μm). Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³ 52
- 4.3 Basic image-based features and their role in organoid phenotype diversity. a-c Uniform Manifold Approximation and Projection (UMAP) of organoid-level features marked by DNA (DAPI) staining intensity (b), actin (Phalloid/FITC) staining intensity (c) and permeability (DeadGreen) staining intensity d Distribution of organoid size for all control (DMSO) treated organoids. e Replicate correlation of organoid size for control treated organoids. f UMAP representation of DMSO treated and drug treated organoids. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³ 53
- 4.4 Experimental batches and their impact on organoid phenotype. a UMAP of organoid level features stratified by organoid line and colored by experimental batch. 54

- 4.5 Organoid phenotype-profiles capture organoid viability. a Representative example images of negative- (0.1% DMSO) and positive control treated organoids ($2.5\mu\text{M}$ bortezomib, cyan = DNA, magenta = actin, yellow = cell permeability; average images were selected and embedded in black background; scale bar: $50\mu\text{m}$). b, Dose-dependent-trajectory of bortezomib treatment effect. UMAP of organoid morphology at different bortezomib doses and (right panel) dose-dependent trajectory for three representative organoid lines. For visual purposes, trajectory inference was limited to partition 1, the left-hand set of measurements within the UMAP, representing ca. 95 % of all imaging data. Panel a created with support from Johannes Betge. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³ 55
- 4.6 Organoid phenotype-profiles capture treatment specific changes in organoid viability. a Distribution of organoid size at different concentrations of Paclitaxel. Shown is a random sample of 30% of all Paclitaxel treated organoids for this and following figures. b Distribution of organoid number per well at different concentrations of Paclitaxel. c Example images of Do₂₂T organoids treated with Paclitaxel. d Dose-response relationship of organoid size and paclitaxel dose. Do₂₂T and Do₄₆T are highlighted. e UMAP of organoid morphology highlighting Do₂₂T organoids treated at different concentrations of Paclitaxel. f UMAP of organoid morphology highlighting Do₄₆T organoids treated at different concentrations of Paclitaxel. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³ 57
- 4.7 Organoid phenotype-profiles reflect ATP-dependent viability measurements. a Association of organoid size of selected example lines with organoid viability determined by luminescence-based, ATP-dependent viability profiling with CellTiter-Glo (CTG), which was performed in parallel with imaging on a subset of drug treatments. b UMAP visualization of dead organoids within our dataset, based on supervised machine learning of organoid viability using classifiers trained on positive- (high-dose bortezomib and SN-38) and negative (DMSO) controls (live-dead classifiers, LDC). c Association of organoid size of selected example lines with organoid viability determined by luminescence-based, ATP-dependent viability profiling with CellTiter-Glo (CTG), which was performed in parallel with imaging on a subset of drug treatments for benchmarking. d UMAP visualization of dead organoids within our dataset, based on supervised machine learning of organoid viability using classifiers trained on positive- (high-dose Bortezomib and SN-38) and negative (DMSO) controls (live-dead classifiers, LDC). e Association of LDC and example organoid features (size, DAPI, actin and permeability dye intensities) with benchmark CTG viability read out. Figure created with support from Jan Sauer (LDC classifier training) and adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³ 58
- 4.8 Distribution of drug activity scores. a Distribution of AUROC scores across all studied organoid models and treatments. b Number of active treatments per organoid line. c Relationship of drug activity and predicted organoid viability. Figure created with support from Jan Sauer (drug activity classifier training, LDC classifier training) and adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³ 59

- 4.15 Factor 1 extended overview. a GSEA of the CRIS C subtype signature over ranked factor 1 gene expression loadings. b IGF2 and H19 locus in the human genome. c-e Association between treatment activity scores and factor 1 scores for three classes of compounds, targeting IGF1R, MEK and EGFR, respectively. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³. 68
- 4.16 Factor 2 overview. a UMAP of observed organoids. Color labels represent the manually determined morphology labels. b Representative images of solid and cystic organoid phenotypes (cyan = DNA, magenta = actin, yellow = cell permeability; average images were selected and embedded in black background; scale bar: 50 μ m). c GSEA of LGR5 gene expression signature over ranked factor 2 gene expression loadings (ranking from high factor 2 to low factor 2 loading). d Distributions of treatment activity score loadings grouped by targets for factor 2. e Relationship of selected treatment activity scores (AUROC) with factor 2 score. Example images curated with support from Johannes Betge. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³. 70
- 4.17 Factor 2 extended overview. a Projection of factor 2 scores for drug-induced phenotypes. Highlighted are drug targets leading to a significant change in projected factor scores across all organoid lines (ANOVA). b Representative images of organoid phenotypes across 5 increasing concentrations of MEK inhibitor treatment and negative control (0.1% DMSO) (cyan = DNA, magenta = actin, yellow = cell permeability; average images were selected and embedded in black background; scale bar: 50 μ m). c Projected dose-dependent changes in factor 2 scores after treatment with the MEK inhibitor Binimetinib. The horizontal black line indicates median factor 2 values of all binimetinib treatment observations. A loess fit with 95% confidence interval (grey background) is provided. d Dose-dependent changes in LGR5 transcript abundance after treatment with the MEK inhibitor trametinib, as assessed by qPCR, data from 3 (Do27T) and 4 (Do19T) independent replicates are presented as mean + s.e.m. * $p < 0.05$, ** $p < 0.005$, NS = not significant, two-sided Student's t test. p values: Do19T: $p = 0.061$ (0.004 μ M), $p = 0.0196$, (0.02 μ M), $p = 0.0187$ (0.1 μ M), $p = 0.024$ (0.5 μ M), $P = 0.0024$ (2.5 μ M), Do27T: $p = 0.0051$ (0.004 μ M), $p = 0.00038$, (0.02 μ M), $p = 0.045$ (0.1 μ M), $p = 0.048$ (0.5 μ M), $P = 0.090$ (2.5 μ M). Example images curated with support from Johannes Betge. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³. 71
- 5.1 Visual abstract of adenoma model profiling project. Mouse colon organoid models were developed, characterized and subjected to image-based small molecule profiling. The resulting data was used to learn a representation space of organoid states, identify state dependent changes in organoid lipid composition and generate hypotheses about small molecules that potentially have the ability to move organoid states. 75

- 5.2 Establishing organoid models of colon adenoma, a Overview of organoid model establishment. Mouse colon organoids were isolated from a transgenic donor animal carrying an inactive conditional oncogenic KrasG12D allele. Homozygous truncation of Apc via CRISPR and activation of the heterozygous KrasG12D allele lead to four different genetically defined organoid models. b In vitro growth factor dependency of adenoma models. Organoids were cultured in complete or modified medium containing combinations of Wnt3A, R-Spondin1-Fc and EGF for 120h and subsequently imaged. Scalebar = 200 μ m. c Oncogenic KrasG12D increases resistance to Egfr inhibition. Organoid ATP levels were measured 4 days after Gefitinib treatment and adjusted for organoid growth rate. Points represent mean of n=2 independent experiments. Error bars represent standard error of mean. d Erk phosphorylation is increased by oncogenic KrasG12D. Organoid models were cultured with or without Wnt3A and R-Spondin1-Fc for 72h and analyzed for protein levels. p, phospho. e Loss of Apc induces transcription of canonical Wnt-signaling target genes. qRT-PCR for Axin2 and Ccnd in the presence or absence of Wnt 3a and R-spondin1-Fc after 120h of culture. Expression levels are normalized to Sdha and Hprt transcript abundance. Bar graphs represent the mean of n=4 independent experiments. Wilcoxon rank sum test 77
- 5.3 Structural validation of organoid colon adenoma models, a Allele-specific PCR products of colon organoid models isolated from a transgenic mouse with a conditional tamoxifen inducible KrasG12D/+ allele. b Amplicon sequencing result of the murine mutation cluster region ortholog for organoids transfected with an Apc targeting sgRNA and Cas9 carrying plasmid. The sequencing results show the presence of 3 different insertion/deletions within the pool of sgRNA treated organoid models. Wildtype sequences are absent within the CRISPR targeted pool, while mutant sequences are absent in the untreated organoid pool. 78
- 5.4 Molecular profiling of organoid adenoma models. a Differential gene expression of adenoma models. Shown are scaled expression values for the top 125 differentially expressed genes for every organoid line. Selected genes are highlighted. All organoids were cultured for 3 days in WENRAS before exposure to ENR for 4 days. Cell number was controlled between experiments. Whole organoid lysates were analyzed. b Transcript abundance data. Shown are the first two principal components of scaled gene expression data. The proportion of variance of each principal component is listed in parenthesis. c Protein abundance data. Shown are the first and third principal component of scaled protein expression data. The proportion of variance of each principal component is listed in parenthesis. d Lipid species abundance data. Shown are the first two principal components of scaled lipid abundance data. The proportion of variance of each principal component is listed in parenthesis. e Loss of Apc leads to increased expression of proliferation and intestinal stem cell associated genes. Shown is a gene set enrichment analysis of differentially expressed genes between Apc mutant and WT organoids. Intestinal gene expression signatures were used according to Merlos-Suarez et al. NES, normalized enrichment score. f Overview of cellular processes in organoid adenoma models. Shown are selected enriched differential gene expression signatures from Reactome and Merlos-Suarez et al. NES, normalized enrichment score. NES > 0 suggests an enriched/ activated biological process. FDR < 0.1. 80
- 5.5 a Representative up and down-regulated transcriptional processes after loss of Apc. Expression signatures were sourced from Reactome and average log₂ fold changes for included transcripts are illustrated. FDR < 0.1. b Representative up and down-regulated transcriptional processes after activation of oncogenic Kras G12D. Expression signatures were sourced from Reactome and average log₂ fold changes for included transcripts are shown. FDR < 0.1. 81

5.6	Image-based profiling of organoid adenoma models. a Overview of experiments. Organoids were isolated from a transgenic mouse model and genetically edited. Organoids were dissociated and evenly seeded in 384-well plates before perturbation with an experimental small molecule library. After treatment, high-throughput fluorescence microscopy was used to capture the morphology of organoids in 16 selected z-layers and 3 channels. 3D imaging data were projected on a 2D plane using a maximum contrast projection. Here, only pixel areas with the largest contrast among the z-axis were retained. Morphological features were computed based on the projection. Untreated organoid morphology, organoid size and treatment activity scores were integrated with transcript expression, protein abundance, lipid abundance and genotype data in a Multi-Omics Factor Analysis (MOFA) model. Figure created with support from Johannes Betge (graphical presentation). b Uniform Manifold Approximation and Projection (UMAP) of organoid-level features for a random 5% sample out of imaged organoids. The identical sample is used for visualizations throughout the figure. Organoid genotype is colorcoded and representative images are displayed (magenta = DNA, cyan = actin, cell permeability = yellow, scale-bar: 200µm). c Graph-based clustering of organoids by morphology with 8 resulting clusters. d Organoid size distribution. Color corresponds to the log-scaled organoid area (dark blue: minimum size, yellow: maximum size).	82
5.7	Treatment and genotype dependent effects on organoid morphology distribution. a UMAP representation of DMSO treated (vehicle) and small molecule treated organoids. b, UMAP embeddings of four organoid genotypes (baseline state = 0.1% DMSO control-treated organoids), grey background consists of randomly sampled organoids.	83
5.8	Genotype dependent effects on organoid morphology. a Unperturbed organoid profiles from adenoma models were aggregated. Shown are representative individual organoids with selected features. Points show the mean phenotype for each independent biological replicate. Representative, interpretable features and their z-scores relative to all single organoid profiles are shown (magenta = DNA, cyan = actin, cell permeability = yellow, scale-bar: 25µm)	84

- 5.13 Small molecule Wnt signaling inhibitors. a AUROC activity score for three small molecule inhibitors of canonical Wnt signaling. b Target proteins for small molecules within the canonical Wnt signaling cascade with their relative position to the destruction complex (highlighted in blue). c Distributions of lipid abundance loadings grouped by lipid species for factor 1 (Abbreviations: HexCer - Hexosylceramide; PA - Phosphatidate; Cer - Ceramide; SM - Sphingomyelin; LPC - Lysophosphatidylcholine; PS - Phosphatidylserine; PE - Phosphatidylethanolamine; PI - Phosphatidylinositol; PG - Phosphatidylglycerol; DAG - Diacylglycerol; PC - Phosphatidylcholine; TAG - Triacylglycerol; Hex2Cer - Dihexosylceramide; CE - Cholersterolester) 91
- 5.14 Projection of factor 1 scores for treatment-induced phenotypes and viability changes. Highlighted are compounds leading to a significant change in projected factor scores across all organoid lines (ANOVA). Organoid viability is predicted using a random-forest based classifier (LDC) with scores from 0 (no toxicity) to 1 (complete toxicity) 92
- 5.15 GSK₃ beta inhibition dependent morphology in colon organoid models. a Small molecule inhibition of GSK₃ beta (CHIR98014) leads to phenocopying of Apc^{-/-} genotype organoid models. b Shift of morphological features of wildtype and KrasG12D^{+/+} organoid models treated with CHIR98014. Shown is an increase in organoid size (Area) and DNA intensity. c Excerpt of clustering from figure 5.9 d, labeled with known binding activity of listed small molecules. Rucaparib is not member of the cluster and shown for comparison. 93
- 5.16 Factor 2, KrasG12D induced senescence. a Distributions of treatment activity loadings grouped by drug target for factor 2. b Relationship of representative drugs' activity with factor 2 score. Shown are compounds from highlighted groups in panel (a). c Gene set enrichment results of a senescence signature by ian et al. over ranked factor 2 gene expression loadings (ranking from high factor 2 loading to low factor 2 loading, NES = normalized enrichment score). d Distributions of lipid abundance loadings grouped by lipid species for factor 2 (Abbreviations: HexCer - Hexosylceramide; PA - Phosphatidate; Cer - Ceramide; SM - Sphingomyelin; LPC - Lysophosphatidylcholine; PS - Phosphatidylserine; PE - Phosphatidylethanolamine; PI - Phosphatidylinositol; PG - Phosphatidylglycerol; DAG - Diacylglycerol; PC - Phosphatidylcholine; TAG - Triacylglycerol; Hex2Cer - Dihexosylceramide; CE - Cholersterolester). e Relationship of acyl chain length with factor 2 loading. Shown are lipids from highlighted species in panel (d) 95
- 5.17 Factor 3, KrasG12D effects in the context of Apc loss of function. a Distributions of treatment activity loadings grouped by drug target for factor 3. b Relationship of representative mTOR inhibitor activity with factor 3 score. c Gene set enrichment results of a Reactome mTORC1 activation signature over ranked factor 3 gene expression loadings (ranking from high factor 3 loading to low factor 3 loading, NES = normalized enrichment score). d Visual summary of Myc gene set enrichment results for organoid state transitions. Myc signatures were significantly enriched in Apc^{-/-} models and depleted in models with isolated KrasG12D^{+/+} mutation. 97
- 5.18 Projected changes in factor 1 and 2 scores after small molecule treatment a Shows are treatments leading to change in projected factor scores across all organoid lines (ANOVA). Color coding represents predictions based on LDC viability classifier. The structural formula of Ellagic Acid is depicted in the top right. b Organoid treatment with Ellagic acid. Depicted are representative example images for each line (blue = DNA, red = actin, green = cell permeability, scale bar=200µm). 99

Nothing in Biology Makes Sense Except in the Light of Evolution

Theodosius Dobzhansky

1

Introduction

1.1 DISCLOSURE

Parts of this introduction, especially the section on Wnt signaling, have been adapted from own publications, including *Wnt signaling in cancer*³

Details matter, it's worth waiting to get it right.

Steve Jobs

2

Materials and Methods

2.1 PATIENTS

All patients were identified at the University Hospital Mannheim, Mannheim, Germany. We included untreated patients with a new diagnosis of colon or rectal cancer in this study and obtained biopsies from their primary tumors and adjacent normal tissue via forceps based endoscopy. Exclusion criteria were active HIV, HBV or HCV infections. Biopsies were transported in phosphate buffered saline (PBS) on ice for subsequent organoid extraction. Clinical data, tumor characteristics and molecular tumor data were pseudonymized. The study was approved by the Medical Ethics Committee II of the Medical Faculty Mannheim, Heidelberg University (Reference no. 2014-633N-MA and 2016-607N-MA). All patients gave

The beginning is the most important part of the work.

Plato, The Republic

3

Image-based profiling of organoids

3.1 DISCLOSURE

Significant parts of this chapter have been adapted from own manuscripts, including *The drug-induced phenotypic landscape of colorectal cancer organoids*³. The maximum contrast projection method, organoid segmentation method, feature extraction procedure and organoid viability classification (LDC) were previously developed by Jan Sauer as part of his dissertation^{noa}. Image-based profiling experiments were supported by Johannes Betge.

3.2 ESTABLISHING PATIENT DERIVED ORGANOIDS FOR IMAGE-BASED PROFILING

Patient derived organoids can be established from diverse healthy or malignant tissues and have been shown to represent their tissue of origin with respect to morphological and molecular features including gene expression and somatic mutations^{8,35,30,24,4}. To generate personalized cancer models for image-based profiling, I designed and implemented a standardized laboratory workflow to generate patient derived organoids from colorectal cancer samples via endoscopic biopsy (Figure 3.1a). Briefly, fresh patient samples were washed, digested and embedded in a basal membrane extract, a proprietary mixture of extracellular matrix proteins especially rich in Laminin and Collagen 4. The isolated tumor cells were then overlaid with a growth factor rich medium, containing Epidermal Growth Factor (EGF), the BMP-signaling antagonist Noggin and the small-molecule inhibitor A83-01, which inhibits TGF-beta-signaling by targeting the Activin receptor-like kinase family.

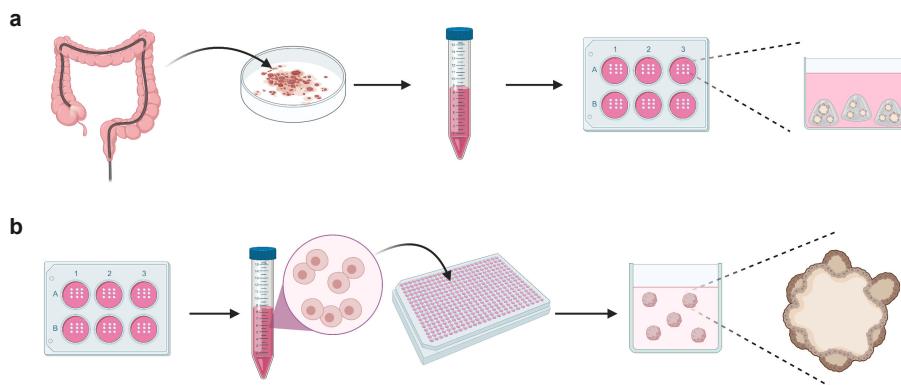


Figure 3.1: Core liquid handling methods a Organoid isolation procedure. Colorectal cancer tissue biopsies were collected via endoscopy, enzymatically removed from extracellular matrix proteins, washed and resuspended in basal membrane extract hydrogel. After solidification of hydrogel domes, organoids were overlayed with growth factor rich culture medium. b Organoid high-throughput experimentation. Colorectal cancer organoids were harvested, partially digested, seeded in hydrogel-coated 384-well plates

Following this protocol, patient derived organoids from 13 patients with colorectal cancer were prospectively developed. Donors to the biobank represented different UICC stages (Figure 5.6a).

Amplicon sequencing of frequently altered genes in colorectal cancer showed molecular profiles characteristic for the disease (Figure 5.6c). Similar to sequencing studies of primary tumors, patient derived organoids harbored a high frequency of APC (6/13), KRAS (8/13) and TP53 (5/13) mutations¹⁹.

On a gene expression level, patient derived organoids mainly represented the canonical consensus molecular subtype CMS 2 of colorectal cancer¹⁰ (Figure 5.6b). No patient derived organoid line with a MSI-high phenotype and the associated CMS 1 molecular subtype was established. Also, no organoid line matched the molecular subtype CMS 4, which is associated with stromal infiltration and TGF β -signaling. These results are in line with previous observations^{30,28} and demonstrate limitations of the organoid culture system, which selects for growth of epithelial cells (favored by canonical Wnt signaling high, BMP4 signaling low) over mesenchymal cells (favored by canonical Wnt signaling low, BMP4 signaling high) *ex vivo*²⁴.

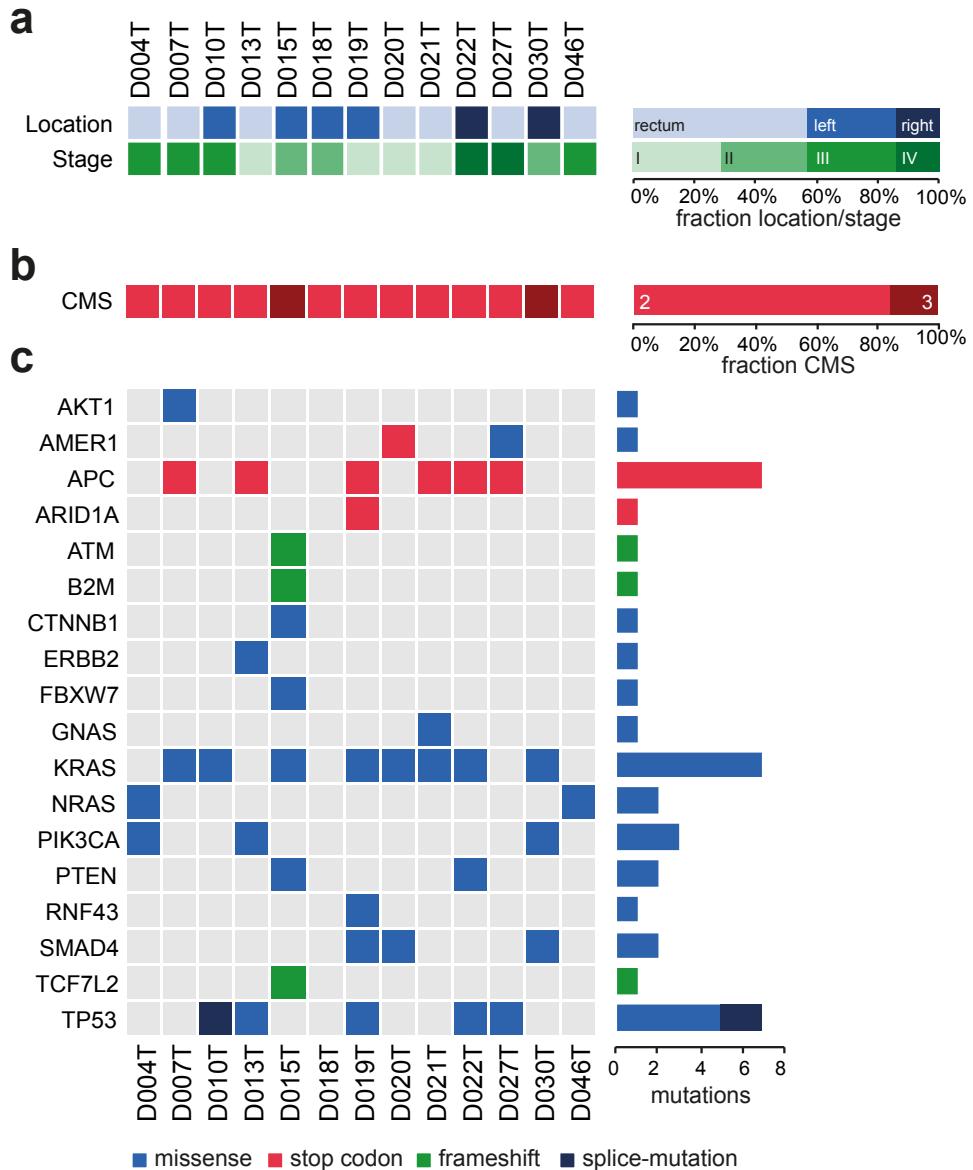


Figure 3.2: Organoid cohort overview a Tumor location (right/left/rectum) and AJCC/UICC stage of colorectal cancers that patient derived organoids were derived from. b Consensus molecular subtypes of organoids determined by RNA expression analysis. c Mutation status in PDOs, as analyzed by amplicon sequencing. Figure created with support from Johannes Betge (graphical presentation), Erica Valentini (sequencing data analysis) and Benedikt Rauscher (CMS type inference). Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³

3.3 ENABLING METHODS FOR HIGH-THROUGHPUT IMAGE-BASED PROFILING OF ORGANOIDS

To systematically measure organoid morphology, I established a platform for high-throughput image-based profiling experiments (Figure 5.8a). The three engineering problems that had to be solved were (1) control of organoid size and density in a 384-well plate format, (2) control of organoid location within the hydrogel for efficient microscopy and (3) maintenance of organoid integrity during automated fixation and permeabilization.

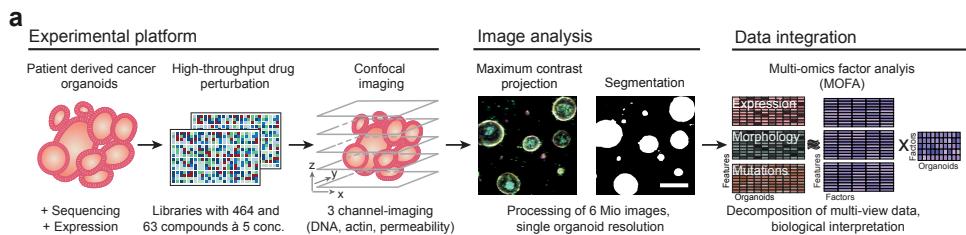


Figure 3.3: Overview of experiments a Organoids were isolated from endoscopic biopsies from patients with colorectal cancer. Organoids were dissociated and evenly seeded in 384-well plates before perturbation with an experimental (464 compounds) and a clinical compound library (63 compounds à 5 concentrations, 842 perturbations across both libraries). After treatment, high-throughput fluorescence microscopy was used to capture the morphology of organoids. The multi-channel (DNA, beta-actin, cell permeability) 3D imaging data was projected, segmented, and descriptive features were extracted to quantify potential drug-induced phenotypes. Untreated organoid morphology, organoid size and drug activity scores were integrated with mRNA expression and mutation data in a Multi-Omics Factor Analysis (MOFA) to increase interpretability of organoid variation. Figure created with support from Johannes Betge (graphical presentation) and adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*.³

A standard protocol for cell-based assays, including image-based profiling, is seeding cells into microwell plates at a fixed cell number. In order to determine cell number, adherent cells are dissociated and counted using optical methods. Patient derived organoids, however, demonstrated a low rate of organoid outgrowth when passaged by complete organoid dissociation down to the single cell level. To improve organoid outgrowth, the dissociation protocol was stopped early, yielding cell clusters of ca. 1-10 cells. These organoid fragments showed an increased outgrowth rate, which could be further improved by treating cells with 10 µM of Rho-Kinase inhibitor Y-27632 (data not shown). To control organoid size and density, organoids were

digested with a modified trypsin derivative, and filtered through a 40 μ m cell strainer to ensure an upper limit of organoid fragment size. To effectively estimate the cell number while maintaining organoid fragments, organoid fragments were titrated based on their ATP concentration, instead of cell count. The ATP concentration of the organoid fragment suspension was determined using an ATP-dependent luminescence readout. After controlling for ATP concentration, the organoid fragment suspension was seeded onto basal membrane extract covered 384 well plates.

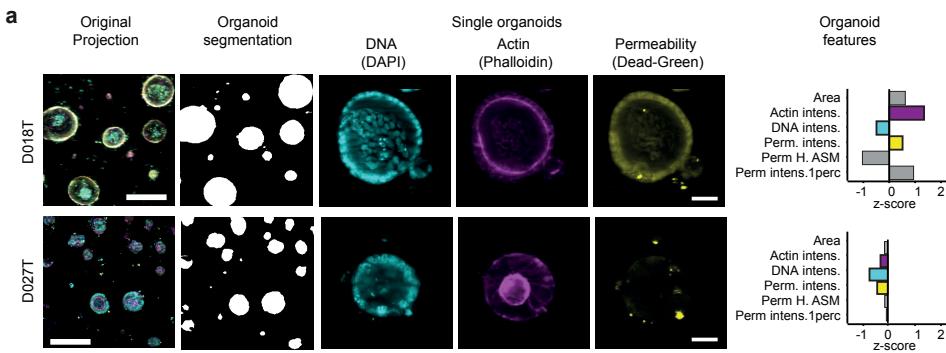


Figure 3.4: Image-based profiling method a The image-processing pipeline illustrated with representative example images from two organoid lines: Organoids were imaged at multiple layers along the z-axis. Images were projected using a maximum contrast projection and segmented using a convolutional neural network, both designed and implemented by Jan Sauer. Descriptive features were extracted from all three channels to quantify phenotypes. Feature plots show the median phenotype of unperturbed organoids, six example features (Area, Phalloidin intensity, DAPI intensity, FITC intensity, FITC Haralick angular second moment (ASM) and FITC intensity 1-percentile) and their z-scores relative to all profiled organoid lines are shown. Figure created with support from Jan Sauer (data processing) and Johannes Betge (graphical presentation). Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*.³

Conventional image-based profiling of adherent cells is based on automatic microscopy of one 2D plane per field of view. Given the 3D growth patterns of organoids, more data has to be acquired to fully capture organoid phenotype. Acquiring multiple planes of imaging data per field of view, however, creates a technical data storage and processing burden. For a fixed 3D volume, the dimensions of the collected data increase linearly with the number of acquired planes and quadratically with the target z-axis resolution. To reduce the observed 3D volume and thus the number of required imaging planes, the vertical distribution of organoid fragments within the basal membrane extract layer was controlled by centrifuging organoid fragments

post-seeding at 500G for 20 minutes at 37 degrees Celsius. The centrifugal force led to an accelerated sedimentation of organoid fragments onto the same optical plane before the polymerization of the hydrogel was complete.

After three days of culture and four days of compound treatment, organoids were fixed and stained for actin (Phalloidin/TRITC), DNA (DAPI), and cell permeability (DeadGreen/FITC) (Figure 3.4a). Subsequently, plates were imaged at multiple z-positions by automated confocal microscopy. During treatment with the hyperosmolar fixative (3% para-formaldehyde in phosphate buffered saline) the protein rich hydrogel underwent an irreversible volume contraction (data not shown). To reduce this artefact, the fixative was supplemented with bovine serum albumin to a final concentration of 1% weight/volume. After image acquisition, 3D data was projected into a 2D plane by applying a maximum contrast projection followed by segmentation with a weakly-supervised convolutional neural network and single-organoid-level feature extraction (Figure 3.4a). In summary, seeding well-quantifiable organoid fragments instead of single cells, centrifuging organoid fragments to reduce the imaged 3D volume, and modifying liquid handling buffers to avoid hydrogel-driven artefacts technically enabled high-throughput image-based profiling of organoid models.

Often when works at a hard question, nothing good is accomplished at the first attack. Then one takes a rest, long or short, and sits down anew to the work. During the first half-hour, as before, nothing is found, and then all of a sudden the decisive idea presents itself to the mind.

Henri Poincare

4

Profiling of organoids identifies molecular determinants of cancer organoid architecture and plasticity

4.1 DISCLOSURE

Significant parts of this chapter have been adapted from own manuscripts, including *The drug-induced phenotypic landscape of colorectal cancer organoids*³. The maximum contrast projection method, organoid

segmentation method, feature extraction procedure and organoid viability classification (LDC) were previously developed by Jan Sauer as part of his dissertation^{noa}. Image-based profiling experiments were supported by Johannes Betge.

4.2 IMAGE-BASED PROFILING CAPTURES THE MORPHOLOGICAL DIVERSITY OF PATIENT-DERIVED CANCER ORGANOIDS

To better understand the diversity of organoid phenotypes and how morphology links to molecular processes, I performed image-based profiling at single organoid resolution with 11 organoid models using compounds targeting developmental pathways (464 compounds), as well as compounds in clinical use (63 compounds in 5 concentrations) together with Johannes Betge (Figure 4.1 a and b). The resulting data comprised morphological profiles for each organoid with 528 phenotypic features that were subsequently reduced into 25 principle components representing 81% of morphological variance.

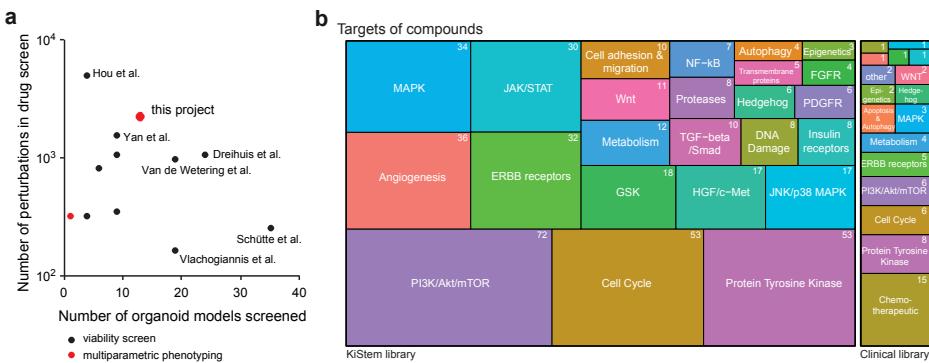


Figure 4.1: Dataset dimensions and compound library overview a Number of organoid models and number of perturbations in previous publications reporting high-throughput drug screenings with patient derived cancer organoids, b Graphical representation of the compound libraries used for drug screening in this project: A library targeting kinases and stem cell pathways (KiStem library, 464 compounds) and a clinical library with 63 drugs in 5 concentrations. Figure created with support from Johannes Betge and adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³

To visualize the heterogeneity of colorectal cancer organoids and drug induced changes across and within cancer organoid lines, the features of ca. 5.5 million profiled organoids were embedded using uniform

manifold approximation and projection (UMAP) (Figure 5.7 a and 4.3 a-c). Most organoid lines showed characteristic bimodal log-normal distributions of organoid size with one component containing small organoids and another component made up of larger organoids with varying, line specific, average size (Figure 5.7 b, and 4.3 d-e). The log-normal-like size distribution likely resulted from intrinsic differences in cellular size and growth rate compounding over time in multicellular organoids.

While DNA and Actin staining intensity were positively correlated with organoid size, cell permeability was negatively correlated and enriched in regions with relatively smaller organoids (Figure 4.3 a-c). Graph-based clustering of this identified 12 regions within the embedding (Figure 5.7 c). When comparing drug-treated organoids to organoids treated with the negative control (DMSO), no clear separation of these two groups, except an increased presence of drug-treated organoids in region 3, was seen. This finding suggested that organoid morphology was distributed on a continuum of phenotypes spanning perturbed and unperturbed conditions of the experiment (Figure 4.3 f).

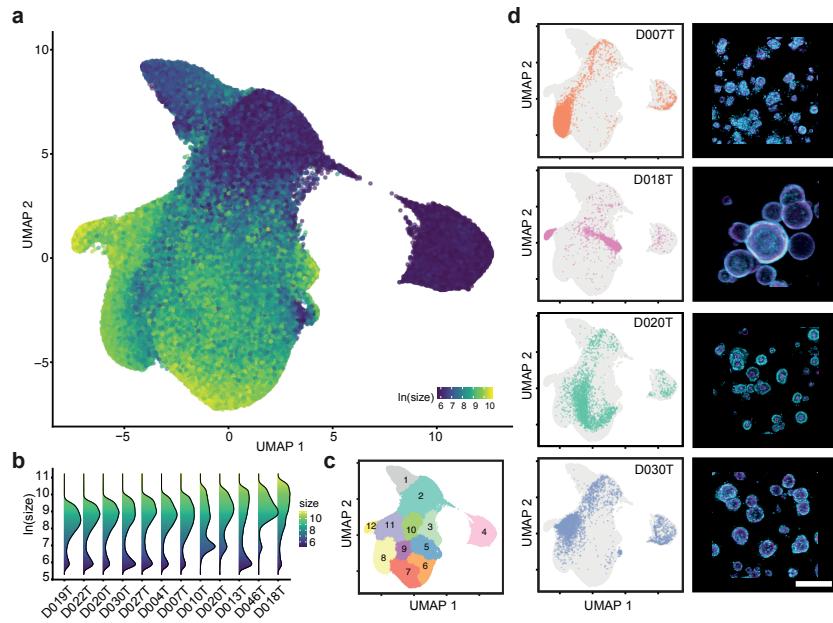


Figure 4.2: Image-based profiling captures the phenotype diversity of patient derived cancer organoids a Uniform Manifold Approximation and Projection (UMAP) of organoid-level features for a random 5% sample out of ca. 5.5 million organoids. The same sample is used for visualizations throughout the figure. Color corresponds to the log-scaled organoid area (dark blue: minimum size, yellow: maximum size). b organoid size distribution across lines. c UMAP representation of DMSO treated and drug treated organoids. Graph-based clustering of organoids by morphology. d UMAP embeddings of selected organoid lines (baseline state / 0.1% DMSO control-treated organoids) representing different morphological subsets, grey background consists of randomly sampled points. Depicted are representative example images for each line (right, cyan = DNA, magenta = Actin, scale-bar: 200µm). Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³

Different organoid lines within the embedding were located in characteristic regions, with organoid size and organoid architecture as primary organizing factors (Figure 5.7 b and d). For example, organoid line Do18T had the largest median organoid size within the dataset and a cystic organoid architecture, while Do20T organoids had a solid architecture and smaller median size. In most cases, organoid lines had two areas of main density, with one of them in regions 2, 3 or 4, reflecting the previously mentioned bimodal size distribution. In summary, image-based profiling of patient derived colorectal cancer organoids showed strong morphological heterogeneity with line dependent differences in size and organoid architecture.

Exploratory data analysis of the relationship between organoid morphology and experimental batch showed

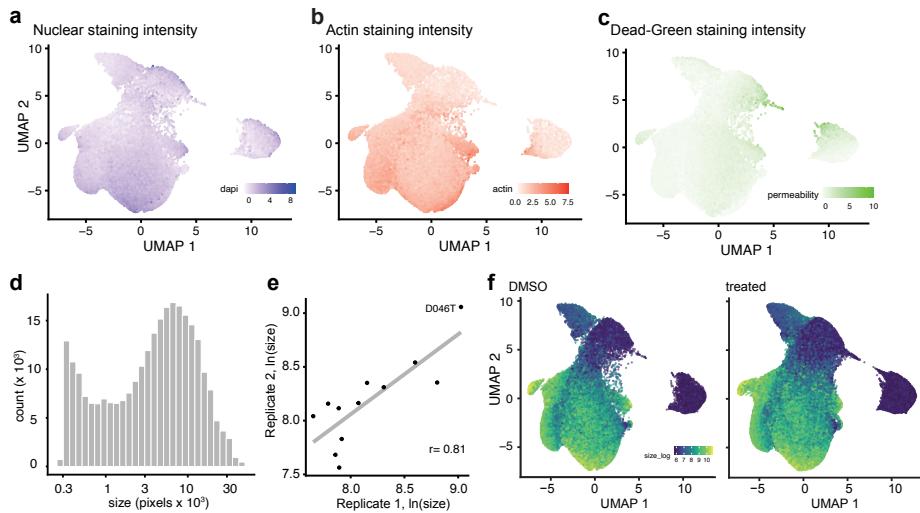


Figure 4.3: Basic image-based features and their role in organoid phenotype diversity. a-c Uniform Manifold Approximation and Projection (UMAP) of organoid-level features marked by DNA (DAPI) staining intensity (b), actin (Phalloid/FITC) staining intensity (c) and permeability (Dead-Green) staining intensity d Distribution of organoid size for all control (DMSO) treated organoids. e Replicate correlation of organoid size for control treated organoids. f UMAP representation of DMSO treated and drug treated organoids. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³

overall reproducible measurements of organoid profiles across experiments (Figure 4.3 e). While objects with a log-area of 8 pixels and larger showed reproducible phenotypes across contexts, smaller objects (mostly dead organoids) showed batch-dependent differences in phenotype.

For example, region 1 within the UMAP embedding was exclusively occupied by observations from batch HC1092-09 and HC1092-10, while region 4 was relatively underoccupied (Figure 4.4 a). Given the confounding of line differences by experimental batches (experimental batches and tested organoid lines were not independent) and the stronger prevalence of batch effects for small objects, no procedure to remove these batch-dependent differences in organoid phenotype were performed.

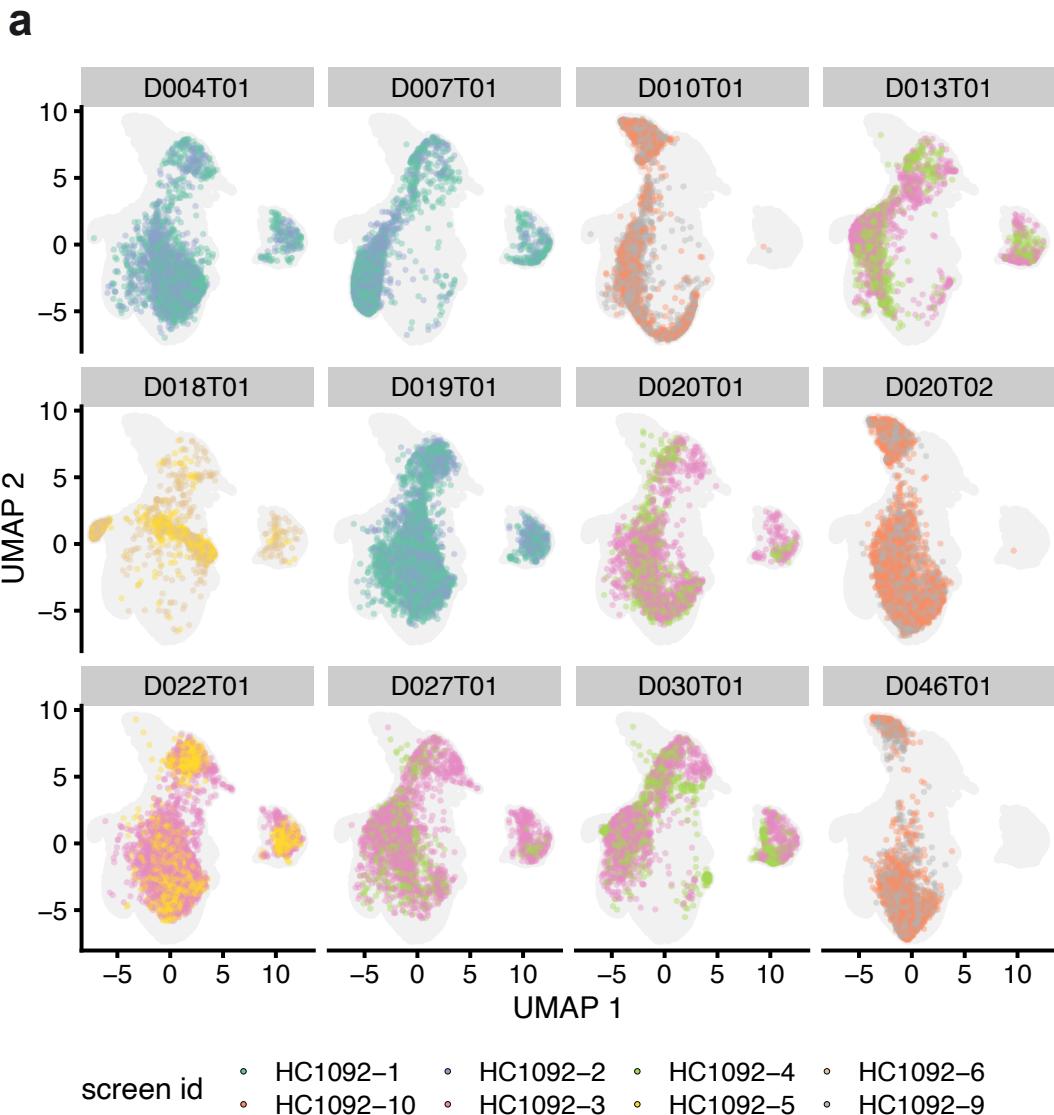


Figure 4.4: Experimental batches and their impact on organoid phenotype. **a** UMAP of organoid level features stratified by organoid line and colored by experimental batch.

4.3 ORGANOID PHENOTYPE-PROFILES CAPTURE ORGANOID VIABILITY

Drug induced changes in cell viability are a basic readout in oncology drug discovery. Prompted by the observation that organoid size was a major factor determining the structure of the phenotype embedding (UMAP and factor 1 in MOFA analysis, see below), I hypothesized that low organoid size was at least partially the result of cell death within the organoid and, more broadly, that phenotype data could be used to estimate organoid viability. Bortezomib, a small molecule proteasome inhibitor with high in-vitro toxicity led to dose dependent organoid death in all organoid lines, thus representing suitable positive controls (Figure 4.5 a). Analogous to pseudotime in single-cell gene expression analysis, dose-dependent trajectories of Bortezomib drug response could be fitted (Figure 4.5 b) using the non-parametric principle curve method. Starting from diverse baseline morphologies, increasing doses of Bortezomib led to a step-wise convergence on a final death-related phenotype, which corresponded to the areas with enrichment of small objects (regions 2, 3 and 4).

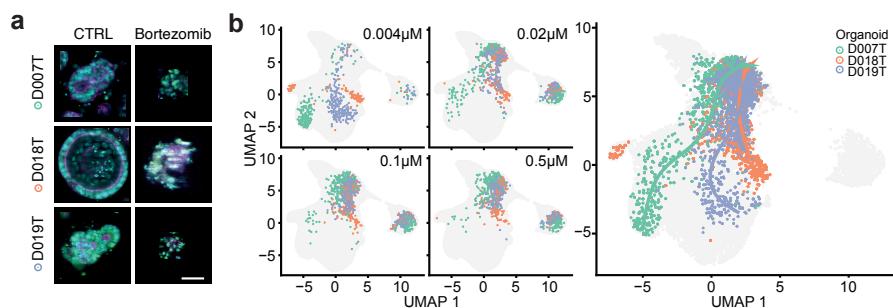


Figure 4.5: Organoid phenotype-profiles capture organoid viability. a Representative example images of negative- (0.1% DMSO) and positive control treated organoids (2.5 μ M bortezomib, cyan = DNA, magenta = actin, yellow = cell permeability; average images were selected and embedded in black background; scale bar: 50 μ m). b, Dose-dependent-trajectory of bortezomib treatment effect. UMAP of organoid morphology at different bortezomib doses and (right panel) dose-dependent trajectory for three representative organoid lines. For visual purposes, trajectory inference was limited to partition 1, the left-hand set of measurements within the UMAP, representing ca. 95 % of all imaging data. Panel a created with support from Johannes Betge. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*.³

Similarly, Paclitaxel, a microtubule disassembly inhibitor, shifted the bimodal size distribution of organoids in

a dose-dependent fashion (Figure 4.6 a), while organoid count remained largely unchanged (Figure 4.6 b). This effect, however, was organoid line-specific, as median organoid size in Paclitaxel sensitive lines (e.g. Do22T) decreased, while the size of other organoids remained unaffected (e.g. Do46T, Figure 4.6 c-f). These observations suggested a link between organoid morphology, especially organoid size, with a loss of cell viability.

To further test the link between organoid morphology and cell viability, I performed a luminescence-based, ATP dependent, cell viability assays (CTG) in parallel with imaging as benchmark. A correlation of CTG viability with organoid size ($r = 0.64$) (Figure 4.6 a) was visible.

To test whether a more accurate prediction of organoid viability was achievable by using all available imaging data, I used a previously trained set of random forest classifiers (live/dead classifiers, LDC). These classifiers were trained on individual organoid phenotype profiles to distinguish between negative and positive control treatments (DMSO, Bortezomib and SN-38).

When applying the classifier to the whole imaging dataset and visualizing predictions via UMAP, organoids within previously identified small-object regions 2, 3 and 4 had the highest probabilities for death (Figure 4.6 b).

Not only organoid size, but also the viability classified (LDC) correlated with luminescence-based viability (CTG) measurements (Figure 4.6 b and c). In fact, the trained classifier, together with organoid size showed the most robust correlation with luminescence-based viability measurements across all profiled organoid lines (Figure 4.6 d), while other simple features, such as DAPI, Actin, and permeability (DeadGreen) intensity were less suitable to predict the viability of organoids.

In conclusion, organoid size is an informative metric to approximate organoid viability, but is biased by line-specific differences organoid size. Models consuming more comprehensive morphological information can achieve even higher predictive performance of organoid viability.

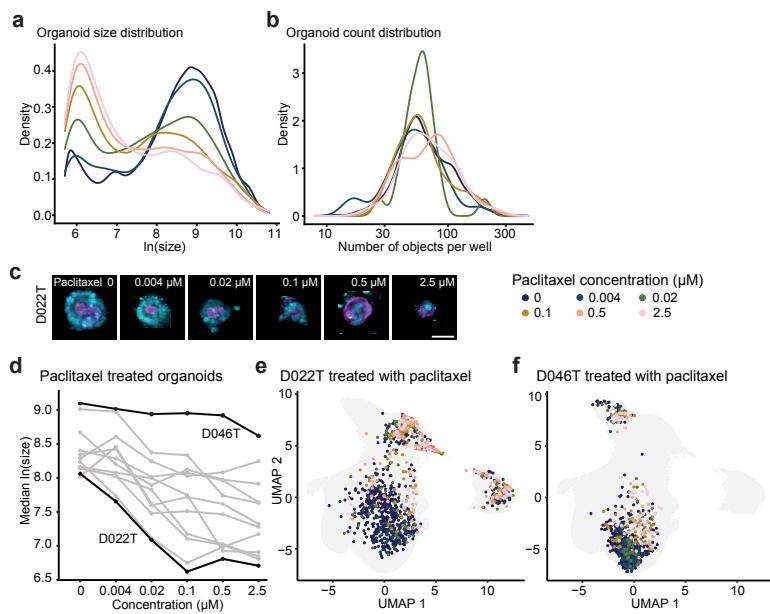


Figure 4.6: Organoid phenotype-profiles capture treatment specific changes in organoid viability. **a** Distribution of organoid size at different concentrations of Paclitaxel. Shown is a random sample of 30% of all Paclitaxel treated organoids for this and following figures. **b** Distribution of organoid number per well at different concentrations of Paclitaxel. **c** Example images of D022T organoids treated with Paclitaxel. **d** Dose-response relationship of organoid size and paclitaxel dose. D022T and D046T are highlighted. **e** UMAP of organoid morphology highlighting D022T organoids treated at different concentrations of Paclitaxel. **f** UMAP of organoid morphology highlighting D046T organoids treated at different concentrations of Paclitaxel. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³

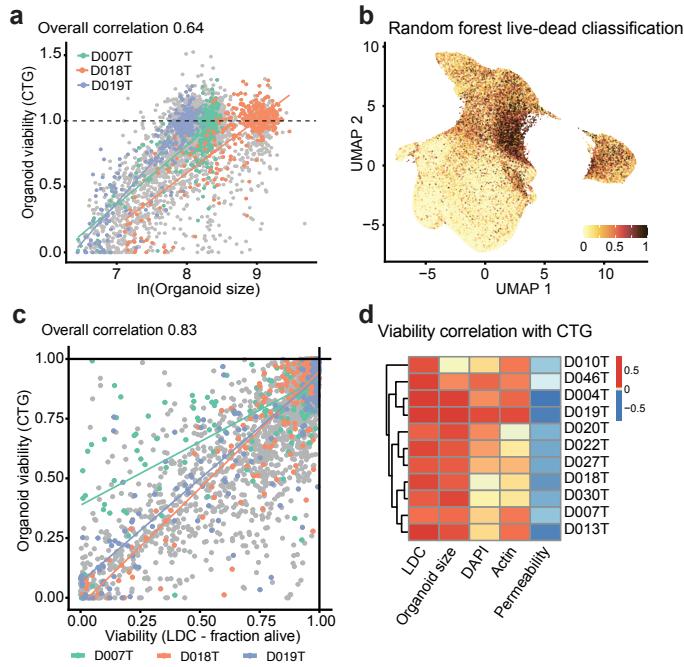


Figure 4.7: Organoid phenotype-profiles reflect ATP-dependent viability measurements. **a** Association of organoid size of selected example lines with organoid viability determined by luminescence-based, ATP-dependent viability profiling with CellTiter-Glo (CTG), which was performed in parallel with imaging on a subset of drug treatments. **b** UMAP visualization of dead organoids within our dataset, based on supervised machine learning of organoid viability using classifiers trained on positive- (high-dose bortezomib and SN-38) and negative (DMSO) controls (live-dead classifiers, LDC). **c** Association of organoid size of selected example lines with organoid viability determined by luminescence-based, ATP-dependent viability profiling with CellTiter-Glo (CTG), which was performed in parallel with imaging on a subset of drug treatments for benchmarking. **d** UMAP visualization of dead organoids within our dataset, based on supervised machine learning of organoid viability using classifiers trained on positive- (high-dose Bortezomib and SN-38) and negative (DMSO) controls (live-dead classifiers, LDC). **e** Association of LDC and example organoid features (size, DAPI, actin and permeability dye intensities) with benchmark CTG viability read out. Figure created with support from Jan Sauer (LDC classifier training) and adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*.³

4.4 DRUG INDUCED ORGANOID PHENOTYPES CORRESPOND TO DRUG MECHANISM OF ACTION

An advantage of image-based phenotyping over cell viability measurements in drug discovery is the ability to use the high dimensional drug-induced phenotype-profiles to identify active but not necessarily lethal drugs and estimate their mechanism of action by unsupervised clustering.

To test whether this approach could be used in cancer organoids, I used a weakly supervised learning approach to identify treatment effect vectors and group them by similarity. With the support of Jan Sauer,

logistic regression models to separate individual compound-treated organoids from unperturbed controls were trained. The resulting normal vector between control- and treated organoid profiles was referred to as the treatment effect vector. Next, every model's ability to separate treated and untreated organoids was scored (AUROC, mean from bootstrapped modeling with values ranging from 0.5 to 1) to identify active treatments that induce a robust change in organoid morphology (Figure 4.8 a, b). I referred to the AUROC as the drug activity score. An AUROC of 1 is associated with a strong difference in organoid morphology, while an AUROC of 0.5 corresponds to no modelable difference. Drug activity (as expressed by an activity score larger than an arbitrary cutoff of 0.85) was necessary but not sufficient for a viability effect (Figure 4.8 c). A fraction of drugs led to identifiable changes in organoid morphology but were not classified as lethal by the LDC model.

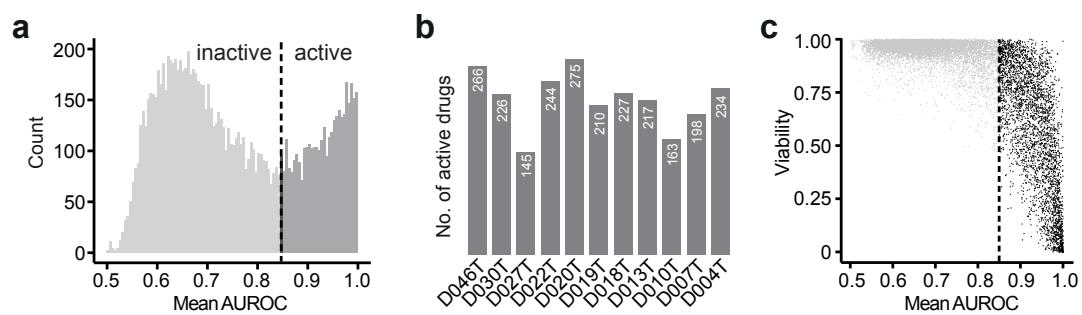


Figure 4.8: Distribution of drug activity scores. a Distribution of AUROC scores across all studied organoid models and treatments. b Number of active treatments per organoid line. c Relationship of drug activity and predicted organoid viability. Figure created with support from Jan Sauer (drug activity classifier training, LDC classifier training) and adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*.³

To test whether active drugs systematically lead to organoid phenotypes that are informative of mechanism of action, the cosine distance between concatenated treatment effect vectors was determined. This approach led to a clustering of specific mode-of-actions, including inhibitors of MEK, Aurora kinase, CDK, mTOR, AKT, EGFR or GSK3 (Figure 4.9 a).

I next validated the previous approach, and tested alternative methods to 1) quantify treatment activity and 2)

cluster treatment effects by similarity. The AUROC score which was previously used to define active treatments, correlated with the euclidean distance of phenotype profiles (Figure 4.10 a) and showed moderate variance when evaluated for stability by bootstrapping (Figure 4.10 b). Similarly, evaluating an alternative clustering method based on phenotype profile averaging (instead of concatenating) followed by Pearson correlation (instead of cosine distance determination) led to an overall similar clustering result of treatments (Figure 4.10 c).

Compounds with related, but not identical, targets also induced related phenotypes, for example MEK inhibitors clustered with specific RAF- and ERK inhibitors (Figure 4.9 c-f) or AKT and PI₃K inhibitors were part of a cluster mainly containing mTOR targeting compounds. The clustering also suggested additional mode-of-actions or off-target effects for well-described compounds. For example, the PKC inhibitor enzastaurin was related to GSK3 inhibitors, substantiating a previously described interaction with the alpha and beta subunits of GSK3^{14,13} (Figure 4.9 f).

To assess whether morphological profiles of active drug treatments were primarily driven by differences in organoid viability, viability predictions were aligned with the phenotypic clustering (Figure 4.9 b). While a large cluster of lethal treatments (including molecules targeting ATM, JAK, PLK, CDK) existed, the majority of clusters were caused by non-lethal phenotypes - including those induced by inhibitors of AKT, mTOR, EGFR or GSK3.

Visual inspection of several phenotypes (Figure 4.9 g) revealed recurring drug target dependent phenotypes. Most notably, MEK inhibitors led to reorganization towards more cystic organoid architecture. These drug target dependent phenotypes were observable across organoid lines and drugs.

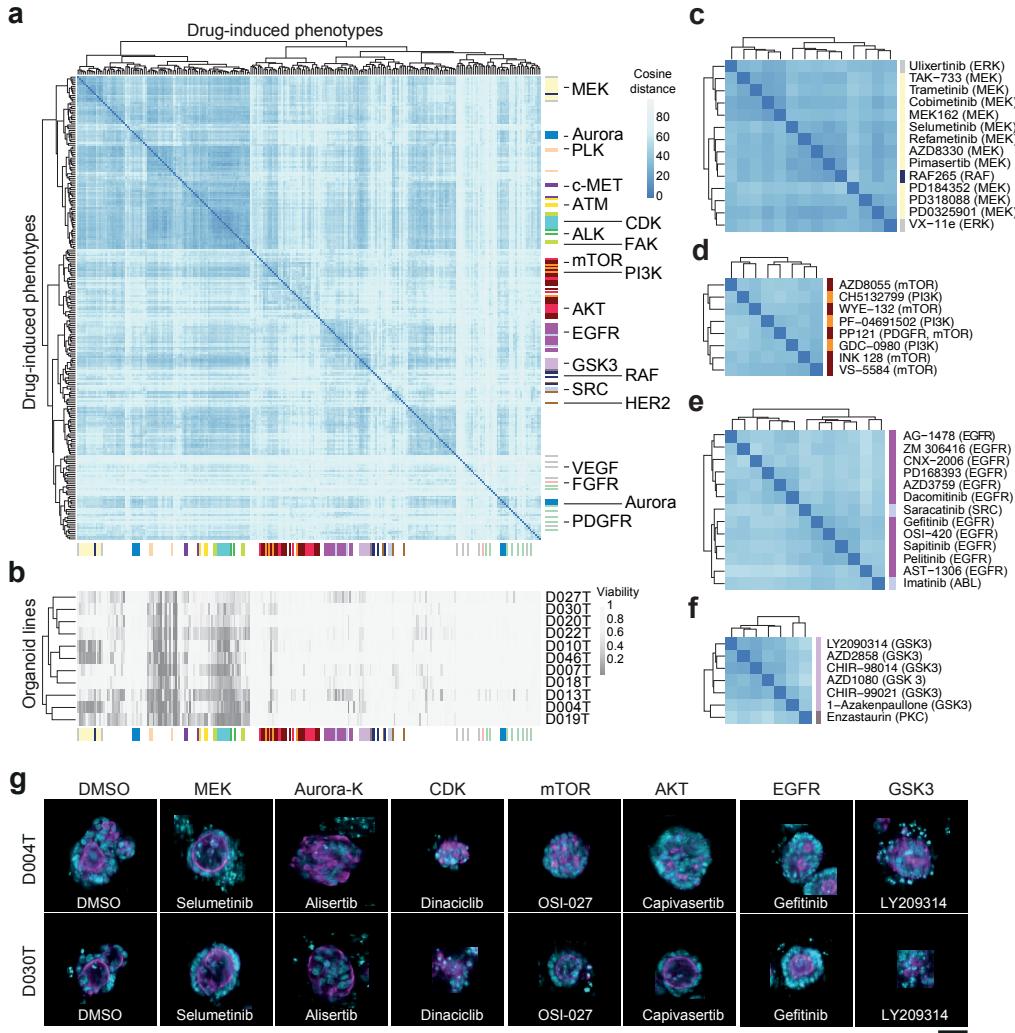


Figure 4.9: Clustering of treatment effects captures mechanism of action. **a** Hierarchical clustering of treatment effect vectors across all observed organoid models. Clustering is based on the cosine distance of concatenated treatment effect vectors. Sidebars are color-coded by the primary annotated target. **b** Viability classifier predictions (LDC classifier) for organoids, arranged according to the clustering in panel a. **c-f** Magnified regions from panel a showing clusters of small molecule inhibitors targeting MEK, mTOR, EGFR, and GSK3. **g** Representative images of treatment-induced organoid phenotypes for seven clusters of small molecules (cyan = DNA, magenta = actin, yellow = cell permeability; average images were selected and embedded in black background; scale bar: 50 μ m). Figure created with support from Jan Sauer (drug activity classifier training, LDC classifier training) as well as Johannes Betge (example picture selection) and adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*.³

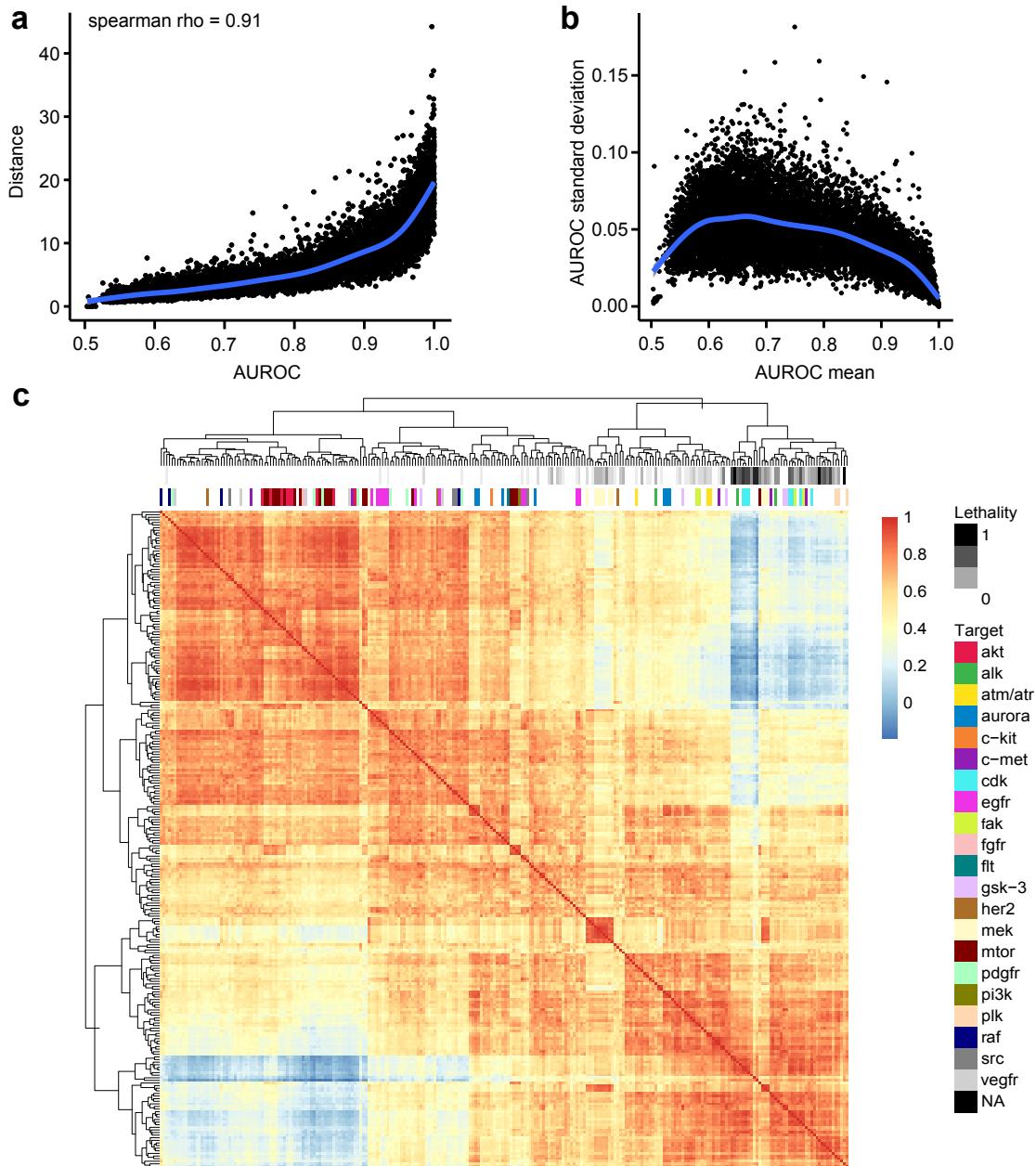


Figure 4.10: Validation of treatment activity and effect quantification. **a** Relationship between euclidean distance of treatment effect vectors and classifier AUROC. The spearman correlation between the two metrics is shown in the top left. Smoothed line represents a loess fit. **b** Relationship between standard deviation of the AUROC score across bootstrapping runs for one treatment and the mean AUROC score of the treatment. Smoothed line represents a loess fit. **c** Hierarchical clustering of treatment effect vectors across all observed organoid models. Clustering is based on the euclidean distance of average treatment effect vectors. Sidebars are color-coded by the primary annotated target.

4.5 MULTI-OMICS FACTOR ANALYSIS IDENTIFIES SHARED FACTORS LINKING MORPHOLOGY, GENOMIC DATA AND DRUG ACTIVITY

A limitation of image-based profiling experiments is that both unperturbed and drug induced phenotypes are challenging to interpret in terms of their underlying biology. I hypothesized that, in the presence of multiple *in vitro* models with both phenotype and genomic measurements, links between the two data modalities can be learned. Based on the observation that organoid morphology representations were distributed in a continuous space, I hypothesized that variation in organoid baseline morphology could be associated with differences in gene expression, as well as drug activity for the 11 cancer organoid lines in our sample.

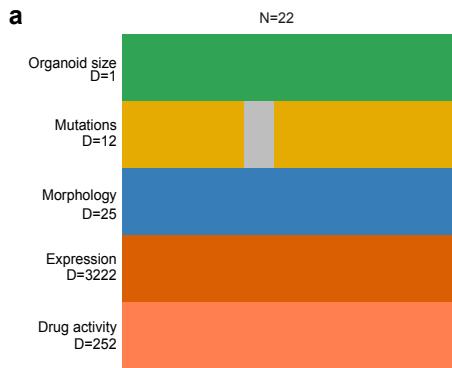


Figure 4.11: Overview of input data for MOFA modeling. 22 observations of 11 models across 5 different views (size, mutation, morphology, gene expression and treatment activity) were integrated in the model. Missing data is shown in grey. "D" represents the vector length of the respective view. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³.

To learn a joint representation of unperturbed organoid morphology, unperturbed organoid size, gene expression, somatic mutations, and drug activity, multi-omics factor analysis (MOFA) was performed. MOFA is a matrix factorization method built upon group factor analysis, with decomposes a set of different measurements into a shared table of factors scoring each observed sample and a set of corresponding tables linking each factor to features in the set of original measurements.

The data used to fit the MOFA model comprised a set of five matrices, each with 22 observations (rows) representing 11 organoid models with 2 replicates per model (Figure 4.13 a). Each matrix represented a data modality/view (mutation, morphology, gene expression, etc.) and had a range of attributes (columns).

When trained with a low number of $k = 3$ factors, MOFA recovered factors explaining ca. 41-24% of variance across the different data modalities, with the first two factors accounting for ca. 29-17% in aggregate (Figure 4.12 a). While gene expression, mutations and drug activity profiles for organoid lines contributed to all factors, factor 1 captured an exceptional amount of variation in median organoid size (ca. 39%). In contrast, factor 2 was primarily capturing variation within baseline organoid morphology (ca. 16%) (Figure 4.12 a).

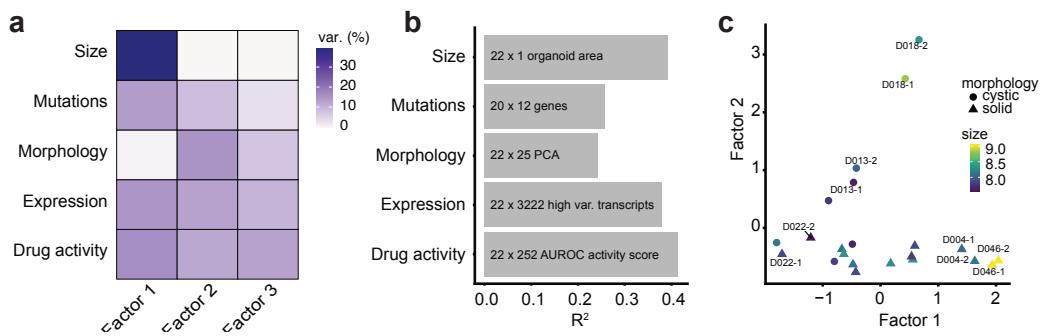


Figure 4.12: Mult-omics factor analysis of organoid profiles. a-b Variance decomposition of the MOFA model. Shown is the variance explained for every factor and data modality, as well as only by modality. c Factor scores for individual observations. The shapes represent the manually determined morphology label and color average organoid size. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³.

Overall, MOFA factors explained up to 40% of variance in median organoid size, drug activity and gene expression, while less than 30% of variance in baseline organoid morphology was explained by the model (Figure 4.12 b). Organoid lines D046T and D004T stood out as lines with the strongest score for factor 1, while lines D018T and D013T had the strongest score in factor 2. Visual inspection of organoids revealed that organoid lines with a higher factor 1 score tended to be larger in size and organoids with high factor 2 score tended to have a more cystic organoid architecture based on manual classification (Figure 4.12 c). No interpretable morphological differences between factor 3 low and high organoids was identifiable, so the subsequent analysis was focused on the first two interpretable factors generated by MOFA.

To ensure that these patterns related to size and morphology were not biased by the dimensionality of views within the model data, I explored the gene expression data directly and plotted organoid models by their manual morpholoical classiication (Figure 4.13 a) and measured size (Figure 4.13 b). Principal component analysis of high variance probes recovered the same observations.

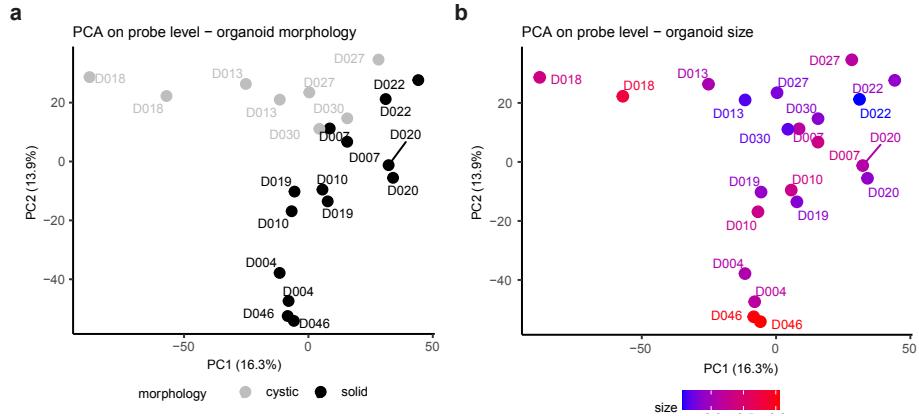


Figure 4.13: Validation of characteristic organoid phenotypes by expression analysis. a PCA of gene expression data for organoid models. Observations are color-coded by the manually determined morphology label. b Identical PCA of expression data, color-coded by average organoid size.

To summarize, MOFA identified factors within the dataset that explained variation between organoid lines across different data modalities, including organoid morphology and median organoid size.

4.6 AN IGF1R SIGNALING PROGRAM IS ASSOCIATED WITH INCREASED ORGANOID SIZE, DECREASED EGFR INHIBITOR ACTIVITY AND CAN BE INDUCED BY mTOR INHIBITION

Differences in organoid size are a contributing factor to intra- and inter-organoid line heterogeneity.

Organoid size was influenced by both organoid line and drug treatments and was associated with factor 1 scores (Figure 4.14 a). An unsupervised gene set enrichment analysis (GSEA) for Reactome pathways across factor 1 loadings showed an enrichment for IGF1R signaling and mitogen-activated protein kinase signaling related genes (Figure 4.14 b).

In fact, the IGF signaling related transcripts H19 (rank 1) and IGF2 (rank 13) were among the strongest contributors to factor 1. This increase in proliferative signaling was confirmed by GSEA of a previously identified intestinal proliferation signature by Merlos-Suarez et al.¹⁸ (Figure 4.14 c).

To better understand clinical correlates to the identified gene expression patterns, I tested for molecular subtypes stemming from an analysis of cancer-cell intrinsic gene expression profiles¹¹. Factor 1 showed an enrichment for CRIS D, a molecular subtype linked to IGF2 overexpressing tumors, loss of IGF2 imprinting and resistance to EGFR inhibitor therapy (Figure 4.14 c). Conversely, I observed a depletion for CRIS C, which has been linked to EGFR dependency (Figure 4.15 a and b). In fact, activity of EGFR inhibitors was the strongest contributor to a negative factor 1 score while IGF1R and MEK inhibitor activity contributed to a positive factor 1 score (Figure 4.14 d-e and Figure 4.15 c-e).

Prompted by the observation that mitogenic signaling, including IGF1R signaling, was underlying factor 1, I hypothesized that other compound treatments could influence the plasticity between the observed organoid states by modulating signaling pathway activity within organoids.

To test whether drug treatments shifted organoid phenotype profiles in factor space, I took advantage of the previous observation that unperturbed and certain perturbed organoids shared similar phenotypic profiles. To this end, the previously estimated factor loading matrix for unperturbed organoid morphology, which was generated during MOFA fitting, was used as a starting point. By generating the pseudoinverse of the loading matrix and multiplying with the average phenotypic profiles of drug-treated organoids, I

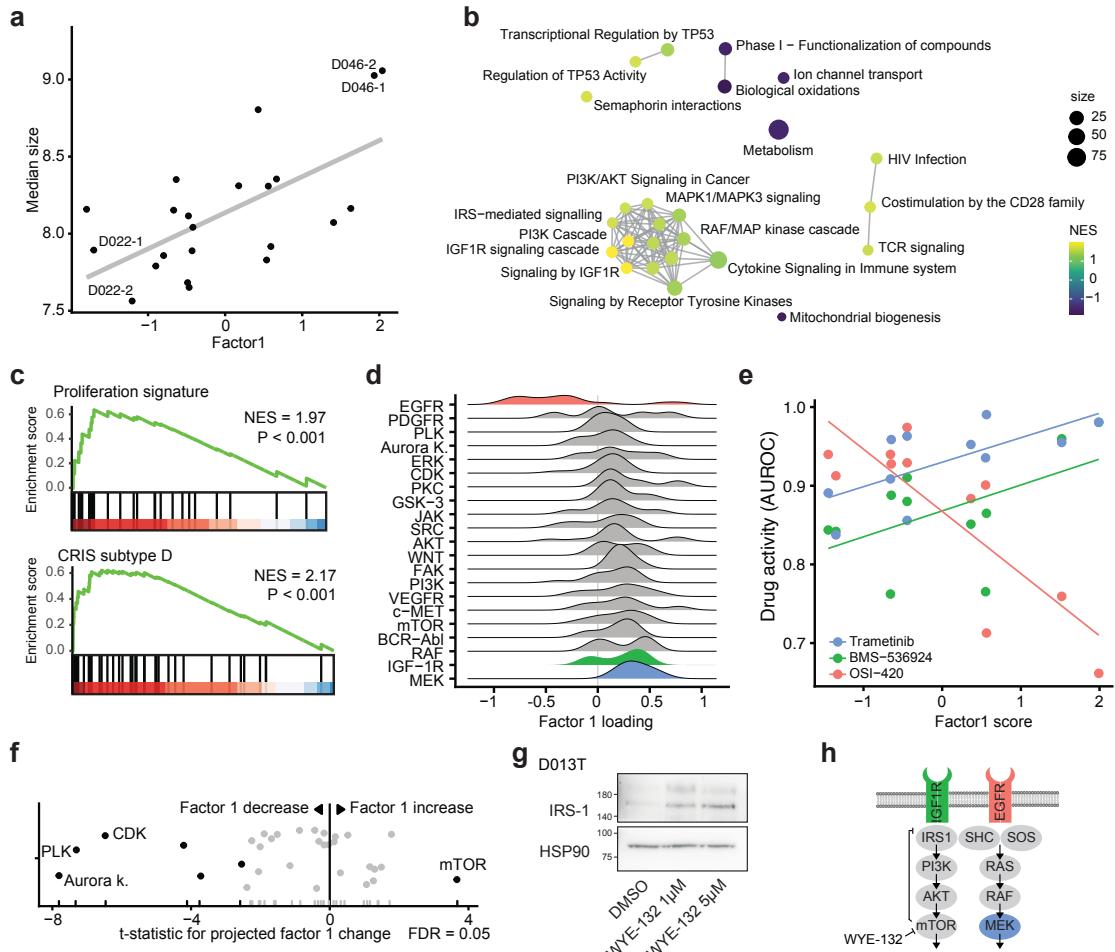


Figure 4.14: Factor 1 overview. **a** Association between organoid size and factor 1 score. **b** GSEA network of factor 1 gene expression loadings. An edge connects Reactome pathways with more than 20% overlap. **c** GSEA of the proliferation signature and the colorectal cancer CRIS-D subtype over ranked factor 1 gene expression loadings (ranking from high factor 1 to low factor 1 loading). **d** Distributions of treatment activity score loadings grouped by targets for factor 1. **e** Relationship of selected treatment activity scores (AUROC) with factor 1 score. **f** Projection of factor 1 scores for drug-induced phenotypes. Highlighted are drug targets leading to a significant change in projected factor 1 scores (ANOVA). **g** Western blot of IRS-1 protein abundance under mTOR inhibition for a representative organoid line (D013T). **h** Illustration of IGF1R signaling pathway with highlighted drug targets. Shown is the disinhibition of mTOR mediated IRS-1 repression after treatment with mTOR inhibitors. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³.

approximated the influence various drug treatments had on biological programs previously identified in unperturbed organoids.

A group of cell cycle related kinase inhibitors targeting polo like kinases, Aurora kinases and cyclin dependent kinases shifted organoids to a low factor 1 score (Figure 4.14 f). In contrast, mTOR inhibitor treatment

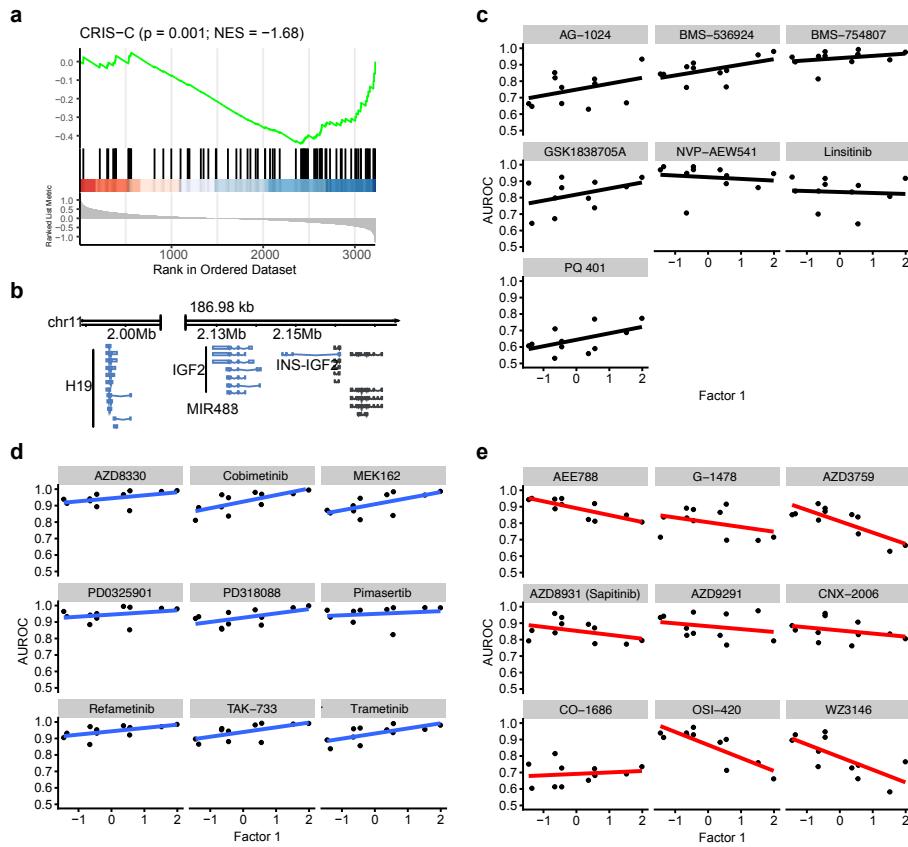


Figure 4.15: Factor 1 extended overview. **a** GSEA of the CRIS C subtype signature over ranked factor 1 gene expression loadings. **b** IGF2 and H19 locus in the human genome. **c-e** Association between treatment activity scores and factor 1 scores for three classes of compounds, targeting IGF1R, MEK and EGFR, respectively. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³.

increased factor 1 scores in cancer organoids (Figure 4.14 f). Given the observation that factor 1 was associated with IGF-1R signaling and mTOR inhibitor treatment led to an increase in factor 1 scores, I hypothesized that mTOR inhibition leads to a reactive upregulation of IGF1R signaling in cancer organoids. In fact, inhibition of mTOR signaling had previously been linked to transcriptional disinhibition of IRS-1 in a negative feedback loop²¹ and reactive induction of IGF1R signaling had previously been described as a resistance mechanism to small molecule mTOR inhibitors in cancer²⁹. When testing this hypothesis in patient derived organoids, a dose-dependent increase of IRS-1 protein abundance in organoids treated with the ATP competitive mTOR

inhibitor WYE-132 was observable (Figure 4.14 g and h).

To summarize, factor 1 described an organoid state with relatively large organoid size, elevated IGF1R dependent mitogenic signaling and relative inactivity of EGFR inhibitor treatment that could be induced by inhibiting an mTOR dependent negative feedback loop in patient derived cancer organoids.

4.7 AN LGR5+ STEMNESS PROGRAM IS ASSOCIATED WITH CYSTIC ORGANOID ARCHITECTURE AND CAN BE INDUCED BY INHIBITION OF MEK

Besides size differences, a particularly strong recurring organoid phenotype was the presence of a cystic organoid architecture, seen for example in untreated D018T organoids and organoids treated with MEK inhibitors (Figure 5.7 and 4.9). In the cystic state, which was observed in factor 2 high organoid lines, organoids consisted of a monolayer of uniform cells lining a central spherical lumen with a distinct apico-basally oriented actin cytoskeleton (Figure 4.16 a and b). This phenotype was reminiscent of organoid morphologies previously seen in APC-/ or Wnt ligand treated human intestinal organoids¹⁷.

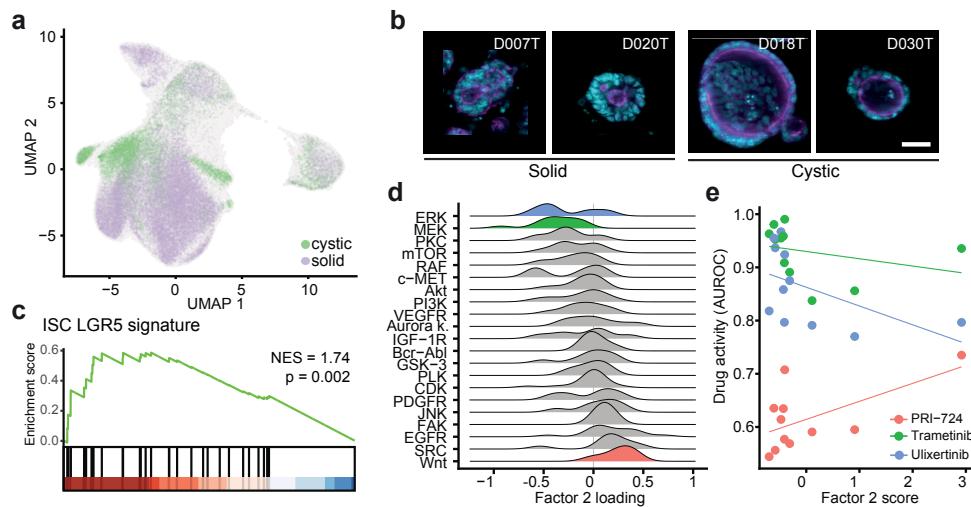


Figure 4.16: Factor 2 overview. a UMAP of observed organoids. Color labels represent the manually determined morphology labels. b Representative images of solid and cystic organoid phenotypes (cyan = DNA, magenta = actin, yellow = cell permeability; average images were selected and embedded in black background; scale bar: 50μm). c GSEA of LGR5 gene expression signature over ranked factor 2 gene expression loadings (ranking from high factor 2 to low factor 2 loading). d Distributions of treatment activity score loadings grouped by targets for factor 2. e Relationship of selected treatment activity scores (AUROC) with factor 2 score. Example images curated with support from Johannes Betge. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³.

To test if factor 2 comprised Wnt signaling and intestinal stem cell identity related gene expression programs, gene set enrichment analyses (GSEA) was performed for cell identity signatures previously by Merlos-Suarez et al.¹⁸. GSEA revealed an enrichment of Lgr5+ stem cell signature-related genes for the factor 2 loadings (FDR=0.002, NES=1.74) (Figure 4.16 c). Next, I wondered whether factor 2 was associated with particular

drug activity or inactivity patterns. Activity of Wnt signaling inhibitors and EGFR inhibitors were the strongest average contributors to a positive factor 2 score (t statistic = 3.02, FDR = 0.046 and t statistic = 3.08, FDR = 0.046, respectively), while activity of ERK and MEK inhibitors were associated with a low factor 2 score (Figure 4.16 d), albeit not significantly. As expected from these results, factor 2 high organoid lines showed a stronger morphological response to the Wnt pathway inhibitor PRI-724. (Figure 4.16 e).

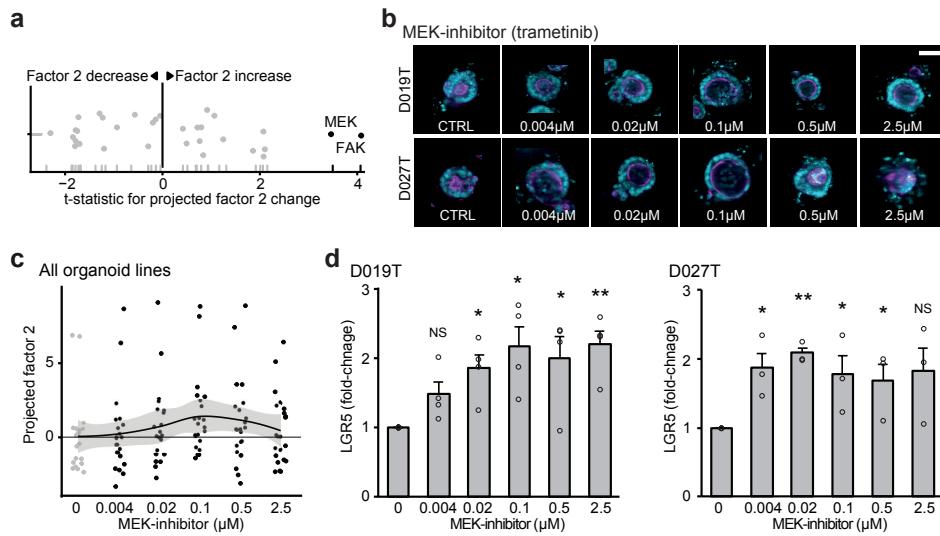


Figure 4.17: Factor 2 extended overview. **a** Projection of factor 2 scores for drug-induced phenotypes. Highlighted are drug targets leading to a significant change in projected factor 2 scores across all organoid lines (ANOVA). **b** Representative images of organoid phenotypes across 5 increasing concentrations of MEK inhibitor treatment and negative control (0.1% DMSO) (cyan = DNA, magenta = actin, yellow = cell permeability; average images were selected and embedded in black background; scale bar: 50 μm). **c** Projected dose-dependent changes in factor 2 scores after treatment with the MEK inhibitor Binimetinib. The horizontal black line indicates median factor 2 values of all binimetinib treatment observations. A loess fit with 95% confidence interval (grey background) is provided. **d** Dose-dependent changes in LGR5 transcript abundance after treatment with the MEK inhibitor trametinib, as assessed by qPCR, data from 3 (D027T) and 4 (D019T) independent replicates are presented as mean + s.e.m. * $p < 0.05$, ** $p < 0.005$, NS = not significant, two-sided Student's t test. p values: D019T: $p = 0.061$ (0.004 μM), $p = 0.0196$, (0.02 μM), $p = 0.0187$ (0.1 μM), $p = 0.024$ (0.5 μM), $P = 0.0024$ (2.5 μM), D027T: $p = 0.0051$ (0.004 μM), $p = 0.00038$, (0.02 μM), $p = 0.045$ (0.1 μM), $p = 0.048$ (0.5 μM), $P = 0.090$ (2.5 μM). Example images curated with support from Johannes Betge. Figure adapted from *The drug-induced phenotypic landscape of colorectal cancer organoids*³.

Next, I again used phenotype profiles of drug treated organoids and approximated how drug treatment shifted organoids along the factor 2 program. I observed MEK and focal adhesion kinase inhibitors significantly shifting all tested organoid lines towards higher factor 2 scores (Figure 4.17 a). This change in

factor 2 scores was concentration dependent for MEK inhibitors and coincided with a visual shift in organoid morphology (Figure 4.17 b and c). Given the observation that factor 2 was enriched for an LGR5+ stem cell signature, the expression of LGR5 transcripts at different concentrations of MEK inhibitor treatment was measured and an analogous dose-dependent increases in transcript abundance was observed (Figure 4.17 d). These findings were in agreement with the observation that MEK inhibitor activity had a negative contribution to factor 2 (4.16 d) - While organoids are shifted to a factor 2 high state by MEK inhibition, within the factor 2 high state itself, organoids are relatively insensitive to this treatment. In summary, factor 2 represents an organoid state with cystic architecture, increased expression of LGR5+ stem cell related genes and increased sensitivity to Wnt signaling inhibitors that could be induced by MEK inhibition.

What I cannot create, I do not understand.

Richard Feynman

5

Profiling to identify revertant therapeutics in pre-malignant models of colon cancer

5.1 MOTIVATION

Based on the observation in the previous chapter that (1) joint representations of organoid morphology and biological state can be learned, and (2) small molecule perturbations can shift organoids in representation space, the following hypothesis can be formulated: If well-characterized small molecules can help explain unknown organoid states, unknown small molecules with desirable properties should be identifiable by their

ability to shift the state of well-characterized organoid models. To test this hypothesis, I generated a set of genetically engineered and well-characterized mouse colon organoid lines that model the initiation of colorectal cancer. I then performed a high-throughput profiling experiment covering FDA-approved, natural and targeted small molecules to identify treatments that moved organoid between states 5.1.

The emergence of colorectal cancer via the chromosomal instability process is a well understood sequence of genetic events that start with hyperactivation of canonical Wnt signaling, often via loss of APC, followed by the hyperactivation of RAS-MAPK signaling, often via oncogenic mutations of KRAS. These two mutations are frequent and significantly co-occurring in colorectal cancer patients²⁷, suggesting an interplay between these two acquired genetic functional events that lead to an evolutionary advantage of premalignant cells. While the genetic events within this process are well understood, no therapeutics targeting the loss of APC or the hyperactivation of KRAS via the frequent G12D activation have been developed for clinical use yet.

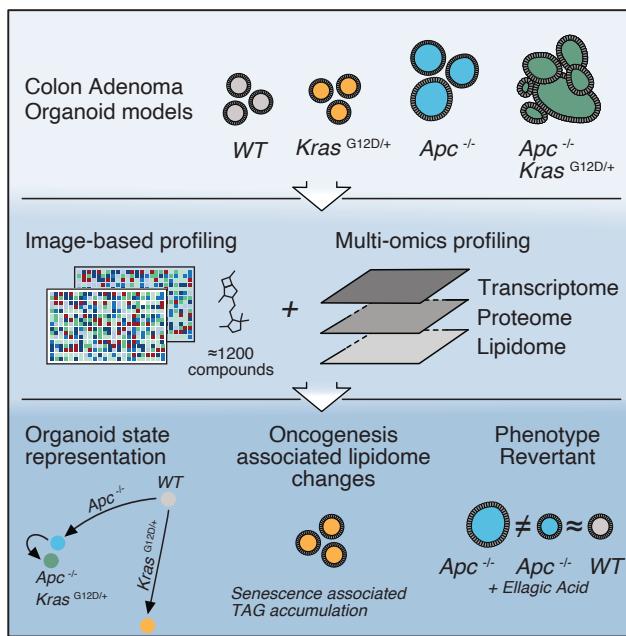


Figure 5.1: Visual abstract of adenoma model profiling project. Mouse colon organoid models were developed, characterized and subjected to image-based small molecule profiling. The resulting data was used to learn a representation space of organoid states, identify state dependent changes in organoid lipid composition and generate hypotheses about small molecules that potentially have the ability to move organoid states.

I genetically engineered mouse colon organoid models carrying *Apc* truncating mutations and/or a *Kras*

G12D allele, thereby modelling the first set of genetic events within the chromosomal instability process leading to colorectal cancer. Using a strategy similar to the approach in the first chapter of this thesis, organoid models were characterized in-depth and subjected to a high-throughput small molecule screen of ca. 1700 FDA-approved substances and experimental compounds. The goal of this project was to (1) understand the biological state changes caused by individual transforming genetic changes in Apc and Kras within colon epithelial cells and (2) identify putative candidates that shift organoid states within the learned representation away from a transformed state.

5.2 GENERATION OF ORGANOID COLON ADENOMA MODELS

APC and KRAS mutant lesions are considered intermediate colon adenomas³. To model the formation of colon adenomas in vitro, I used a transgenic mouse to derive organoid cultures. The transgenic animal carried a conditional tamoxifen inducible KrasG12D/+ allele¹² (Figure 5.2 a). After isolation, I confirmed that extracted colon organoids did not express an activated form of KrasG12D (Figure 5.3a) and defined these organoids as wildtype (WT). To model loss-of-function mutations of the tumor suppressor Apc, the frequently mutated mutation-cluster-region on the APC gene was targeted by CRISPR (Figure 5.2 a). Generated organoids harbored biallelic loss-of-function mutations in Apc (Figure 5.3a). Subsequent activation of oncogenic KrasG12D by treatment with 4-Hydroxytamoxifen led to four distinct organoid adenoma models (Figure 5.2 a and Figure 5.3a-b); wildtype (WT), Apc-/- (A), KrasG12D/+ (K), and Apc-/- / KrasG12D/+ (AK).

Similar to genetically modified human colon organoids^{5,17}, adenoma models showed characteristic niche requirements. Both Apc mutant organoid lines grew independent of the Wnt-signaling activating factors Wnt 3a and R-Spondin1 (Figure 5.2 b) and at an accelerated proliferation rate. In fact, Apc mutant lines showed an increased growth in a Wnt3a and R-Spondin1 free environment when compared to the complete medium. Organoid models with an activated KrasG12D allele were less sensitive to removal of EGF from the media. However, as observed before⁵, the mutant KrasG12D allele was insufficient to compensate completely for the loss of EGF from the medium. Nevertheless, KrasG12D mutant organoid lines were more resistant to

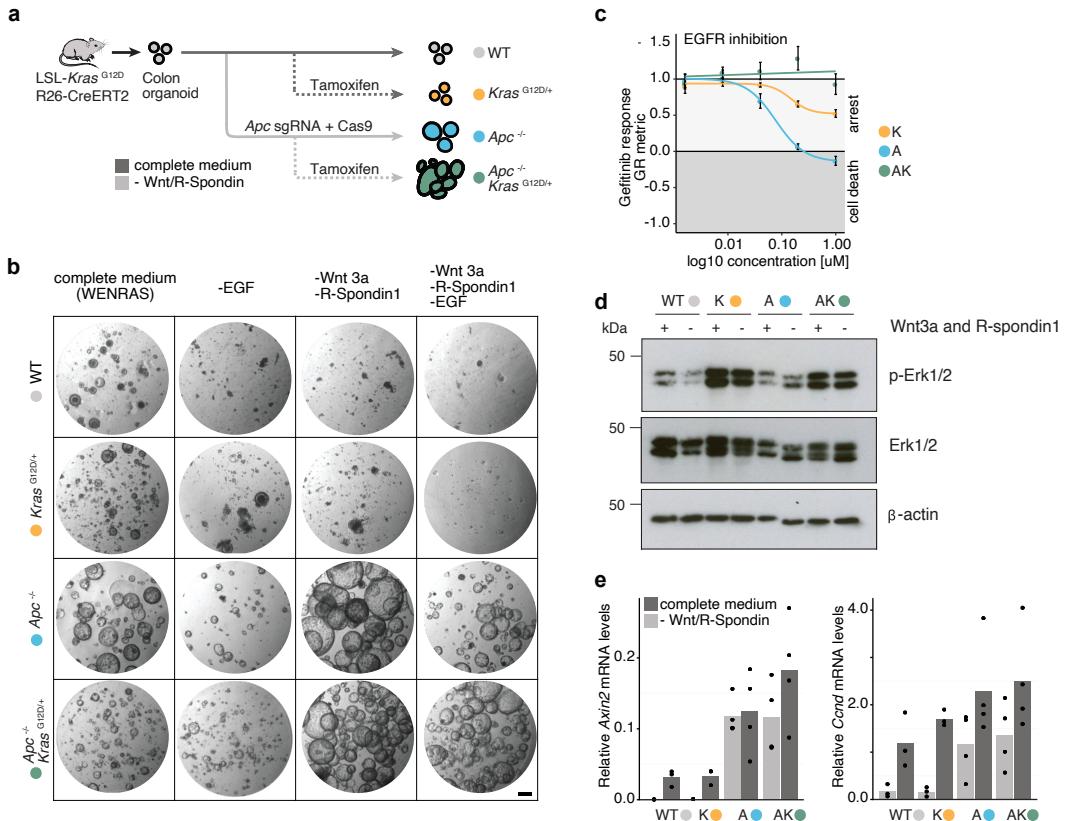


Figure 5.2: Establishing organoid models of colon adenoma. **a** Overview of organoid model establishment. Mouse colon organoids were isolated from a transgenic donor animal carrying an inactive conditional oncogenic KrasG12D allele. Homozygous truncation of Apc via CRISPR and activation of the heterozygous KrasG12D allele lead to four different genetically defined organoid models. **b** In vitro growth factor dependency of adenoma models. Organoids were cultured in complete or modified medium containing combinations of Wnt3A, R-Spondin1-Fc and EGF for 120h and subsequently imaged. Scalebar = 200µm. **c** Oncogenic KrasG12D increases resistance to Egfr inhibition. Organoid ATP levels were measured 4 days after Gefitinib treatment and adjusted for organoid growth rate. Points represent mean of n=2 independent experiments. Error bars represent standard error of mean. **d** Erk phosphorylation is increased by oncogenic KrasG12D. Organoid models were cultured with or without Wnt3A and R-Spondin1-Fc for 72h and analyzed for protein levels. p, phospho. **e** Loss of Apc induces transcription of canonical Wnt-signaling target genes. qRT-PCR for Axin2 and Ccnd in the presence or absence of Wnt 3a and R-spondin1-Fc after 120h of culture. Expression levels are normalized to Sdha and Hprt transcript abundance. Bar graphs represent the mean of n=4 independent experiments. Wilcoxon rank sum test

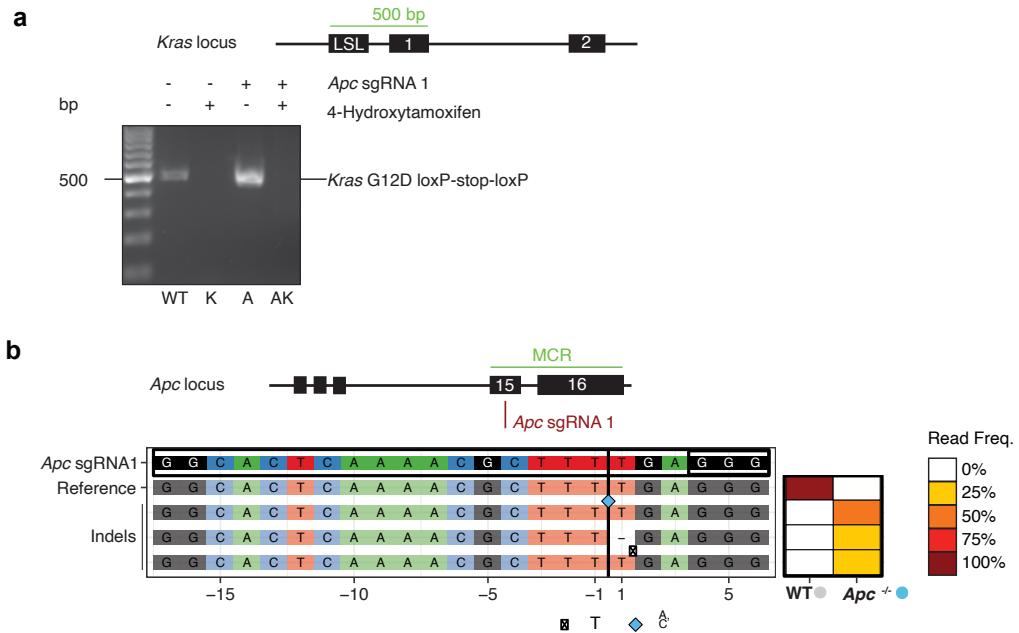


Figure 5.3: Structural validation of organoid colon adenoma models. **a** Allele-specific PCR products of colon organoid models isolated from a transgenic mouse with a conditional tamoxifen inducible KrasG12D/+ allele. **b** Amplicon sequencing result of the murine mutation cluster region ortholog for organoids transfected with an Apc targeting sgRNA and Cas9 carrying plasmid. The sequencing results show the presence of 3 different insertion/deletions within the pool of sgRNA treated organoid models. Wildtype sequences are absent within the CRISPR targeted pool, while mutant sequences are absent in the untreated organoid pool.

pharmacological inhibition of Egfr signaling (Figure 5.2 c). In conclusion, organoid model genotypes were reflected in characteristic growth factor dependencies in experimental conditions.

Next, I investigated the effects of mutations in Apc and Kras on both canonical Wnt- and Erk dependent signaling. While the presence of the KrasG12D/+ allele led to an increase in Erk-phosphorylation across models, Apc-/- / KrasG12D/+ organoids showed no marked additional increase in Erk-phosphorylation when compared to KrasG12D/+ organoids (Figure 5.2 d). Moreover, Apc-/- / KrasG12D/+ adenoma models showed no significant differences in expression of the Wnt target genes Axin2 and Ccnd when compared to Apc-/- single-mutant models (A) ($p > 0.34$ for all conditions, Wilcoxon rank sum test) (Figure 5.2 e). These results indicate that organoid adenoma models show genotype-dependent activity of characteristic signaling pathways, while there is no extensive crosstalk between the Apc-/- and KrasG12D/+ allele in mouse colon

organoids that is directly reflected in canonical Wnt- and Erk dependent signaling.

5.3 MOLECULAR PROFILING OF ORGANOID MODELS

To explore comprehensive molecular differences between organoid models, I next performed transcriptome, proteome and lipidome profiling of all four organoid models (Figure 5.4). Transcriptome profiling of organoid models showed an increased expression of the stem-cell marker Lgr5 and Wnt-signaling regulators such as Nkdr1, Notum, Wif1 and Znrf3 in Apc mutant organoid lines (Figure ?? b). To the contrary, Apc wildtype organoid lines showed an increased expression of epithelial differentiation markers, such as Krt20, Alpp and Abcb1 (P-glycoprotein). Overall, the number of genes with significant expression changes after Apc loss was 2.5 times greater compared to isolated KrasG12D activation (FDR = 0.1, Apc-/-: 44.5%, KrasG12D/+: 18.3% of assessed genes).

A related observation was made during the analysis of protein abundance. Again, Wnt signaling regulators (Axin2, Notum) were enriched in Apc mutant organoid lines and the number of significantly regulated proteins after Apc loss was 2.5 times greater compared to an isolated KrasG12D activation (FDR = 0.1, Apc-/-: 260, KrasG12D/+: 105 assessed proteins).

Principal component analysis of both transcriptome, proteome and lipidome data showed related axes of variation across measurements. In all observed modalities, the first principal component captured differences between Apc wildtype and Apc mutant organoid models, while the second (in case of proteomics measurements the third) principal component captured differences between wildtype and KrasG12D/+ single-mutant models (Figure 5.4b, 5.4c and 5.4d). In every modality, a high degree of similarity was observed among Apc-/- and Apc-/- / KrasG12D/+ organoid lines. While activation of oncogenic KrasG12D in wildtype organoids led to global changes in transcript, protein and lipid expression, these changes were not as pronounced in organoids without functional Apc. In fact, only the mRNA expression of 91 genes was significantly altered between Apc-/- and Apc-/- / KrasG12D/+ organoids (FDR = 0.1).

To explore activated biological processes, gene set enrichment analysis on organoid mRNA expression data was performed. The strongest changes in gene expression after loss of Apc were linked to an increased

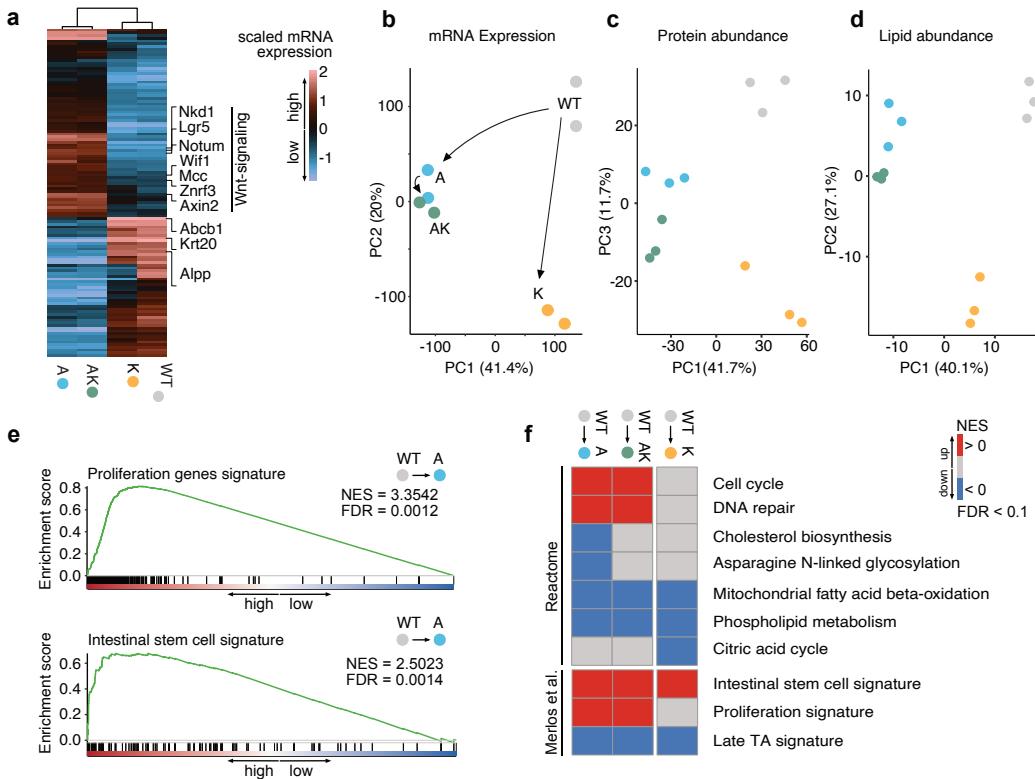


Figure 5.4: Molecular profiling of organoid adenoma models. **a** Differential gene expression of adenoma models. Shown are scaled expression values for the top 125 differentially expressed genes for every organoid line. Selected genes are highlighted. All organoids were cultured for 3 days in WENRAS before exposure to ENR for 4 days. Cell number was controlled between experiments. Whole organoid lysates were analyzed. **b** Transcript abundance data. Shown are the first two principal components of scaled gene expression data. The proportion of variance of each principal component is listed in parenthesis. **c** Protein abundance data. Shown are the first and third principal component of scaled protein expression data. The proportion of variance of each principal component is listed in parenthesis. **d** Lipid species abundance data. Shown are the first two principal components of scaled lipid abundance data. The proportion of variance of each principal component is listed in parenthesis. **e** Loss of Apc leads to increased expression of proliferation and intestinal stem cell associated genes. Shown is a gene set enrichment analysis of differentially expressed genes between Apc mutant and WT organoids. Intestinal gene expression signatures were used according to Merlos-Suarez et al. NES, normalized enrichment score. **f** Overview of cellular processes in organoid adenoma models. Shown are selected enriched differential gene expression signatures from Reactome and Merlos-Suarez et al. NES, normalized enrichment score. NES > 0 suggests an enriched/ activated biological process. FDR < 0.1 .

proliferation rate (Figure ?? e). Gene set enrichment analysis of published intestinal cell-proliferation and stem cell signatures showed an enrichment of both signatures in Apc^{-/-} organoids (Figure ?? e)¹⁸. In contrast, a signature for differentiating, transit-amplifying cells was depleted. Gene set enrichment analysis of Apc^{-/-} / KrasG12D/+ double-mutant organoids showed the same results.

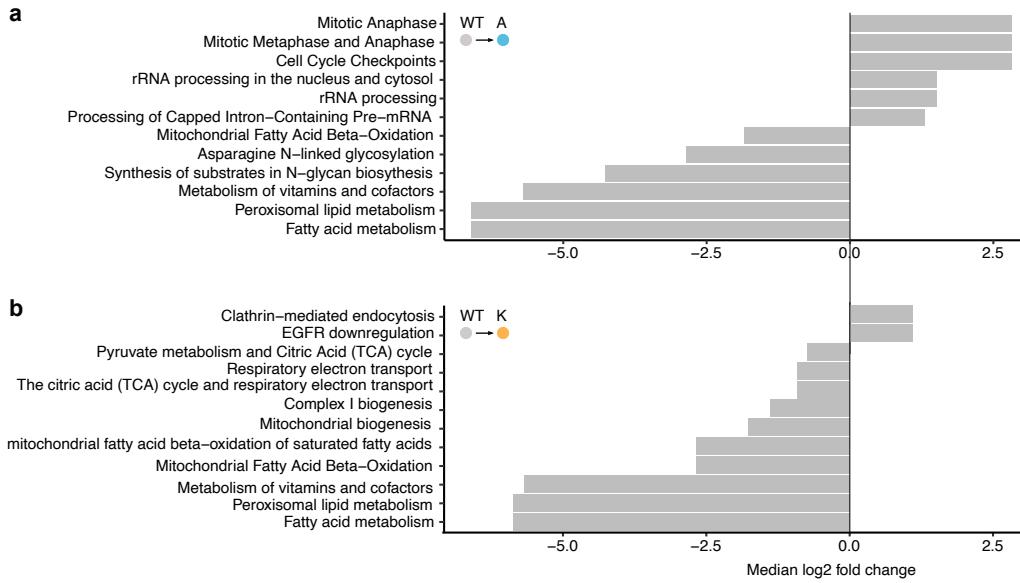


Figure 5.5: **a** Representative up and down-regulated transcriptional processes after loss of Apc. Expression signatures were sourced from Reactome and average log2 fold changes for included transcripts are illustrated. FDR < 0.1. **b** Representative up and down-regulated transcriptional processes after activation of oncogenic Kras G12D. Expression signatures were sourced from Reactome and average log2 fold changes for included transcripts are shown. FDR < 0.1.

Next to these published signatures, I explored the enrichment of curated gene sets from the Reactome database⁹. Here, both Apc^{-/-} and Apc^{-/-} / KrasG12D/+ double-mutant lines showed a positive enrichment of cell cycle and DNA repair related genes when compared to wildtype organoids (Figure 5.5a). Unique to the KrasG12D/+ organoid line was a decreased expression of citric acid cycle and respiratory chain related genes (Figure 5.5b). This effect, was not observed in Apc^{-/-} / KrasG12D/+ double mutant organoids (Figure 5.4f). In addition, organoid models with an KrasG12D/+ genotype showed a downregulation of the EGFR receptor, in line with a potential negative feedback response to hyperactivated RAS-MAPK signaling (Figure

5.5b). Both *Apc*-/- and *KrasG12D*/+ organoid models showed a strong reduction of lipid metabolism and beta-oxidation (Figure 5.5a,b).

In summary, loss of *Apc* leads to a global shift in transcript, protein, and lipid abundance in colon organoids, including a strong increase in cell proliferation associated genes. Activation of isolated oncogenic *KrasG12D* leads to pronounced reduction in citric acid cycle related gene expression while this phenotype was not seen in organoid models with an additional loss of *Apc*.

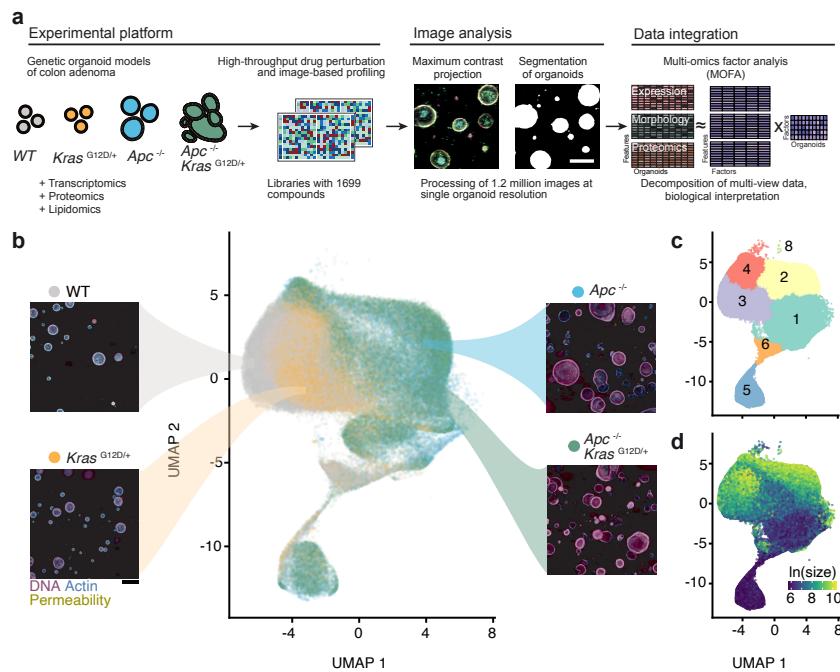


Figure 5.6: Image-based profiling of organoid adenoma models. **a** Overview of experiments. Organoids were isolated from a transgenic mouse model and genetically edited. Organoids were dissociated and evenly seeded in 384-well plates before perturbation with an experimental small molecule library. After treatment, high-throughput fluorescence microscopy was used to capture the morphology of organoids in 16 selected z-layers and 3 channels. 3D imaging data were projected on a 2D plane using a maximum contrast projection. Here, only pixel areas with the largest contrast among the z-axis were retained. Morphological features were computed based on the projection. Untreated organoid morphology, organoid size and treatment activity scores were integrated with transcript expression, protein abundance, lipid abundance and genotype data in a Multi-Omics Factor Analysis (MOFA) model. Figure created with support from Johannes Betge (graphical presentation). **b** Uniform Manifold Approximation and Projection (UMAP) of organoid-level features for a random 5% sample out of imaged organoids. The identical sample is used for visualizations throughout the figure. Organoid genotype is colorcoded and representative images are displayed (magenta = DNA, cyan = actin, cell permeability = yellow, scale-bar: 200 μ m). **c** Graph-based clustering of organoids by morphology with 8 resulting clusters. **d** Organoid size distribution. Color corresponds to the log-scaled organoid area (dark blue: minimum size, yellow: maximum size).

5.4 IMAGE-BASED PROFILING OF ORGANOID MODELS

To measure how organoids change their biological state as a response to small molecule perturbation, I used the previously developed image-based profiling method to observe organoid morphology. Organoid models of four different genotypes were perturbed with a library of ca. 1700 compounds and morphological profiles were computed (Figure 5.6 a). A UMAP projection of the first 25 principal components representing single-organoid morphology showed distinct genotype-dependent morphological states for viable organoids (Figure 5.6 b). Graph based clustering of organoid morphology profiles resulted in 8 different clusters (Figure 5.6 c). While developed organoids within cluster 4 and 3 were enriched for *Apc*+/+ organoid models, cluster 2 and 1 were populated by *Apc*-/- models. Analogous to gene expression, lipidomics and proteomics representation space, *Apc* mutant organoid models were less distinct from each other than organoids with a WT and isolated *KrasG12D*/+ genotype (Figure 5.7 b). While developed organoids that present with a larger organoid area showed distinct genotype-specific morphologies, small and dead organoids clustered together across genotypes within cluster 5 (Figure 5.6 c and d).

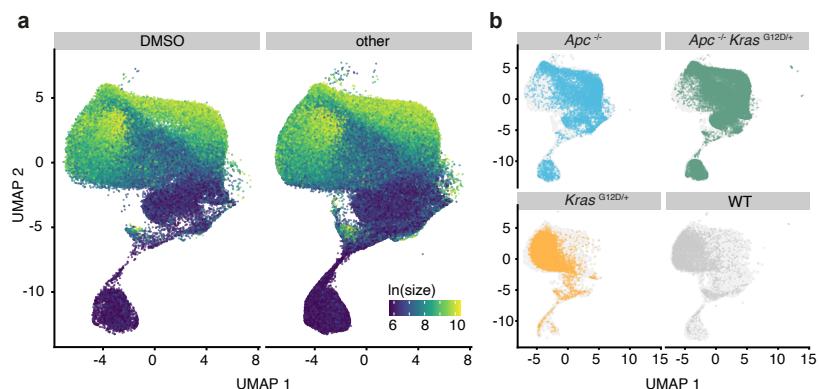


Figure 5.7: Treatment and genotype dependent effects on organoid morphology distribution. a UMAP representation of DMSO treated (vehicle) and small molecule treated organoids. b, UMAP embeddings of four organoid genotypes (baseline state = 0.1% DMSO control-treated organoids), grey background consists of randomly sampled organoids.

The distribution of DMSO-treated organoids and small molecule perturbed organoids in morphological space overlapped strongly (Figure 5.7 a), possibly because of large amount of treatments that did not alter organoid morphology.

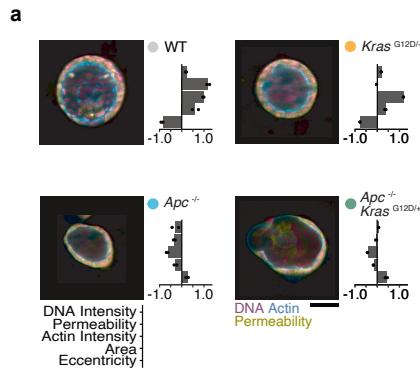


Figure 5.8: Genotype dependent effects on organoid morphology. a Unperturbed organoid profiles from adenoma models were aggregated. Shown are representative individual organoids with selected features. Points show the mean phenotype for each independent biological replicate. Representative, interpretable features and their z-scores relative to all single organoid profiles are shown (magenta = DNA, cyan = actin, cell permeability = yellow, scale-bar: 25 μ m)

When comparing the morphologies of different organoid models in detail, characteristic differences were identifiable (Figure 5.8 a). DMSO-treated *Apc*+/+ organoids showed a strong, regular apical actin cytoskeleton (high average actin intensity) that organized the multicellular formation into a regular-patterned spherical morphology (low average eccentricity). In contrast, *Apc*-/- organoids showed a relative lack of a regular actin cytoskeleton (low average actin intensity) and a irregular, non-spherical morphology (high average eccentricity).

In summary, organoid models showed genotype-dependent differences in morphology. Analogous to differences in molecular state, a primary source of variation was caused by loss of the tumor suppressor gene *Apc*. Organoids with truncated *Apc* presented with a higher proliferation rate, increased overall DNA staining intensity and loss of the spherical apical actin cytoskeleton that was observed in *Apc*+/+ organoid models.

5.5 QUANTIFYING SMALL MOLECULE INDUCED PHENOTYPES ACROSS ORGANOID MODELS

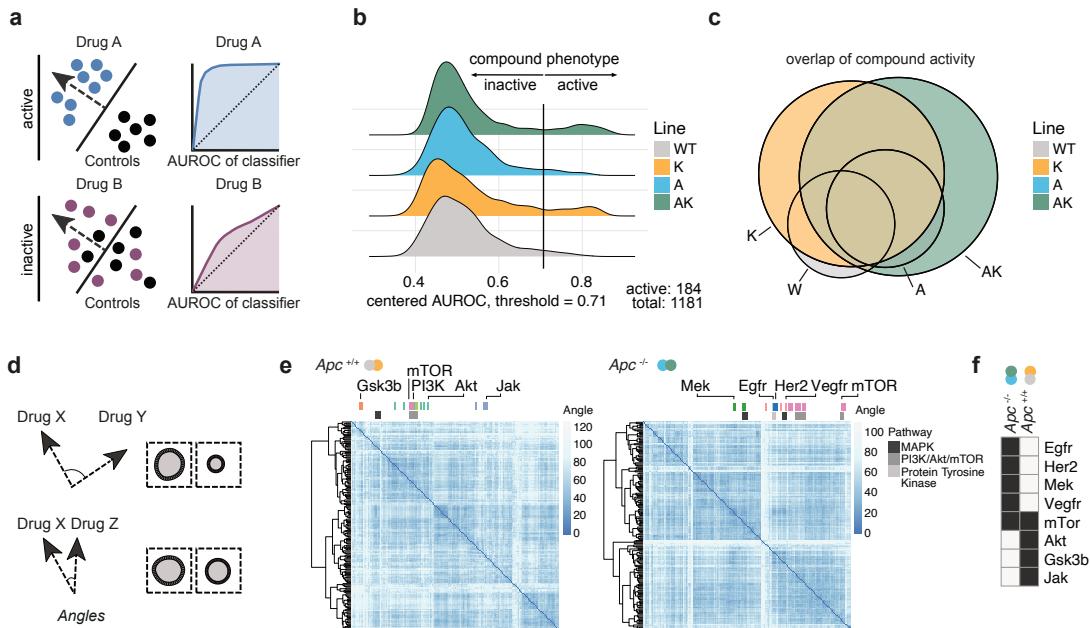


Figure 5.9: Treatment activity scoring. **a** A logistic regression classifier is trained to distinguish morphology profiles of individual treated and untreated organoids across all available replicates. Afterwards, the classifier is applied to a validation set of organoids and the classification performance is estimated using the area under the receiver operating characteristic curve (AUROC) metric. Method implemented by Jan Sauer. **b** Distribution of treatment activity scores for all organoid lines, replicates and perturbations. AUROC scores were centered around 0.5 and treatments for this particular analysis were termed active when classification accuracy exceeded three times the Median Absolute Deviation (MAD) of the AUROC score distribution, an arbitrary threshold. A set of 184 compounds (16% of all screened small molecules) met the activity criteria. **c** Overlap of active compound treatments across organoid models. Shown is an Euler diagram of active compounds for each line. Apc^{-/-} (abbreviated A), double mutant (Apc^{-/-} - KrasG12D^{+/+}, abbreviated AK), Kras G12D (abbreviated K), and wildtype (abbreviated WT) are color coded. Plot diagnostics: diagError: 0.011, stress: 0.002. **d** Identifying related treatment phenotypes. Normal vectors of treatment specific classifiers were compared by calculating the angular distance (related to cosine similarity, ranging from 0-180 degrees). Small angular distance between vectors correspond to a high similarity between the treatment-induced organoid phenotypes. Method implemented by Jan Sauer. **e** A map of compound induced phenotypes for Apc mutant and Apc wildtype organoids. Highlighted are clusters of compound induced phenotypes with related targets. Normal vectors for Apc mutant and Apc wildtype organoids were concatenated before angular distance calculation. Method implemented by Jan Sauer. **f** Treatment induced phenotypes by organoid genotype. Shown are significantly enriched treatment induced phenotypes for Apc mutant and wildtype organoid models. Clusters of similar phenotypes were tested for overrepresentation of known molecular targets using Fisher's exact test. Significantly enriched targets are shown. Method implemented by Jan Sauer.

To study the effect that small molecule perturbations had on organoid models, the classification based approach developed during the study of human cancer organoid phenotypes in the previous chapter was used. Briefly, for every treatment and genotype, a logistic regression classifier was trained to distinguish

DMSO-treated organoids from treated organoids. The classification performance, expressed as the AUROC, was used to determine the activity of a treatment. A high AUROC score (approaching 1) is observed for compounds that lead to a treatment-induced organoid morphology that is very distinct from DMSO treated organoids. In contrast a low AUROC (minimum of 0.5) is observed for compounds where the model's classification performance approaches random guessing (Figure 5.9 a).

Related to the approach chosen in the previous chapter, active treatments were identified based on the AUROC score that a classifier reached. Given differences in the distribution of AUROC scores between lines, with KRAS G12D/+ organoid lines being shifted towards higher AUROC values, I centered the distribution of AUROC scores around 0.5 and defined an arbitrary activity threshold at 3 times the Median Adjusted Deviation for all tested models (Figure 5.9 b). Most active compounds were shared among Apc+/+ and Apc/- models. As seen across other data modalities, the state of the Apc allele was a primary source of variation (Figure 5.9 c). However, the higher classification performance seen in organoids with the Kras G12D/+ allele, led to a larger number of active drugs identified for these models. A possible reason for this systematic difference in the number of active treatments might be a larger number of organoids seen in the images of these models. This difference might be linked to the previously described increased colony-forming capacity seen in Kras G12D/+ colon cancer models²².

Next to identifying differences across treatment activity, I was interested in the question whether morphological treatment effects for compounds varied by organoid genotype. Given the observation that the primary source of variation for treatment activity was the state of the Apc allele, I aggregated organoid lines by their Apc allele for further analysis. Similar to the approach taken in the previous chapter, normal vectors of the logistic regression classifiers were compared by estimating the enclosed angle using the cosine distance (Figure 5.9 d). Normal vectors for organoids with the same Apc allele were concatenated. The resulting clustering of normal vectors by their similarity showed an enrichment for small molecules with related mechanism of action (Figure 5.9 e and f). For example, EGFR inhibitors were significantly enriched in Apc-mutant organoid lines, while GSK3B-inhibitors, which lead to a stimulation of canonical Wnt signaling, were only enriched in Apc-wildtype organoid models.

To summarise the section above, organoid models showed genotype-specific treatment activity patterns. For

example, GSK₃B-inhibitors were active in Apc-wildtype organoids and showed a characteristic treatment-induced phenotype in these models.

5.6 MULTI-OMICS FACTOR ANALYSIS IDENTIFIES SHARED FACTORS LINKING FUNCTIONAL AND STRUCTURAL BIOLOGICAL VIEWS

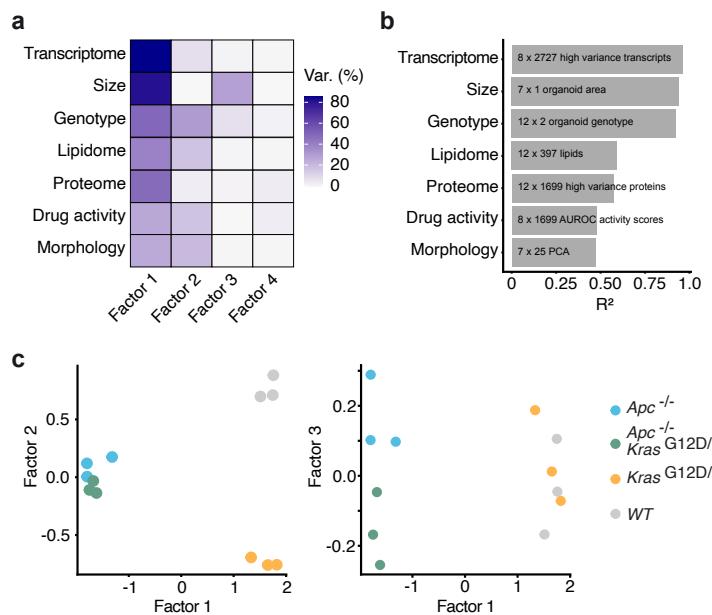


Figure 5.10: Multi-omics factor analysis to identify shared factors linking morphology, size, gene expression, lipidomics, proteomics, genotype and treatment activity. **a**, Percent variance explained by the MOFA model for each factor. Untreated organoid morphology, organoid size and treatment activity scores were integrated with genotype, proteomics, lipidomics and mRNA expression data. **b**, Cumulative proportion of total variance explained by each experimental data modality within the MOFA model. **c**, Visualization of samples in factor space showing factors 1 and 2 as well as factor 1 and 3. Shown are independent replicates for each organoid line.

To comprehensively model the biological state of organoid models and explore potential interventions that

move organoids in state-space, I performed multi-omics factor analysis (MOFA). Analogous to the process described in the previous chapter, feature matrices from different sources were processed and factorized using $k=4$ factors (Figure 5.10 a and 5.14 a). The learned model was based on both functional (e.g. treatment activity) and structural (e.g. genotype, proteomics, lipidomics and mRNA expression) information. To reduce the dimensionality of data modalities with a high amount of features, only high variance features from gene expression and proteomics analysis were used.

The resulting factorization explained ca. 90% (gene expression) to 50% (morphology) of variance across the analyzed views, of which the first three factors captured the majority of variance, Figure 5.10 a). The learned model explained most variance within the mRNA expression and genotype data, while measurements within the organoid morphology data had the lowest explained variance (Figure 5.10 b). Visual inspection of factors as well as exploration of factor loadings within the genotype view showed that factor 1 explained state differences caused by Apc loss of function, while factor 2 explained state differences caused by the activation of KrasG12D in an Apc+/+ genotype (Figure 5.10 c and 5.14 b). In contrast to factor 2, factor 3 captured differences between Kras+/+ and KrasG12D/+ organoids with Apc loss of function, albeit with low overall explained variance.

While the initial number of factors is a user-defined feature within MOFA, the method automatically drops excess factors if they are not considered effective based on an applied automatic relevance determination (ARD) prior. Increasing the number of factors above $k=4$ in this analysis, did not lead to an increased number of interpretable factors. In fact, factor 4 already did not capture differences between organoid genotypes and was not interpretable from a biological point of view (Figure 5.14 b). This observation and the fact that this study explored the biological effects of two oncogenic genetic events both in isolation and in concert reveals, I speculate, a conceptual intersection of representation learning and the theory of genetic interactions which will not be further explored in this thesis.

5.7 A CANONICAL WNT SIGNALING ASSOCIATED PROGRAM CAUSED BY APC LOSS

To understand the molecular changes associated with factor 1, factor loadings for mRNA expression data were analyzed using Reactome gene-set enrichment analysis (Figure 5.12 a). Three clusters of biological processes

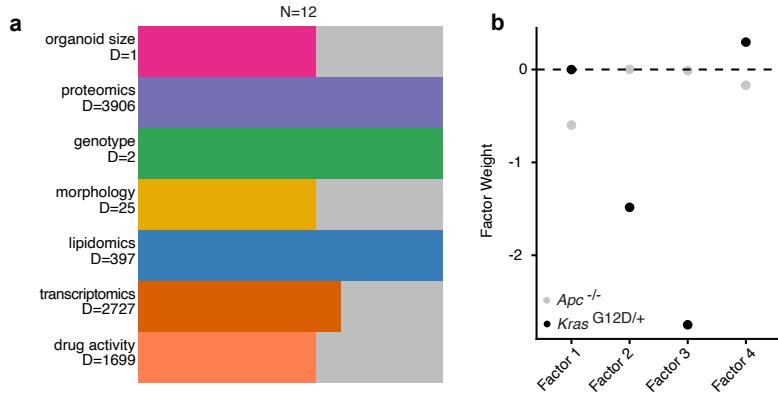


Figure 5.11: Multi-omics factor analysis input data and loadings. **a** measurement modalities, dimensionality and number of measurements. A third replicate of measurements were available for proteomics and lipidomics only. **b** Factor loadings for genotype information.

were significantly associated with a negative factor loading, caused by *Apc* loss-of-function: 1) Mitotic Anaphase related processes, including spindle checkpoints; 2) Mitotic S-phase, including DNA replication and 3) DNA repair mechanisms, including homology directed repair.

In line with the enrichment of processes seen in cell proliferation, factor 1 loadings were associated with an enrichment of a previously described intestinal proliferation signature (Figure 5.12 c) and an LGR5+ intestinal stem cell identity signature (Figure 5.12 b). These findings are in line with the long-standing evidence that loss of *Apc* leads to a hyperactivation of canonical Wnt signaling, which in turn leads to increased intestinal cell proliferation and Myc-dependent changes towards a stem-like cell state^{23,26}.

When focusing on compound activity related factor loadings, a low factor 1 score was significantly linked to increased sensitivity towards small molecules known to target microtubuli and focal adhesion kinase (FAK, Figure 5.12 d). This morphological sensitivity was presented itself primarily as reduced organoid size and number relative to the DMSO vehicle control (Figure 5.12 e).

The average treatment activity scores of small molecules targeting Wnt signaling was associated with increased factor 1 scores (Figure 5.12 d), while microtubuli and FAK inhibitors were associated with decreased factor 1

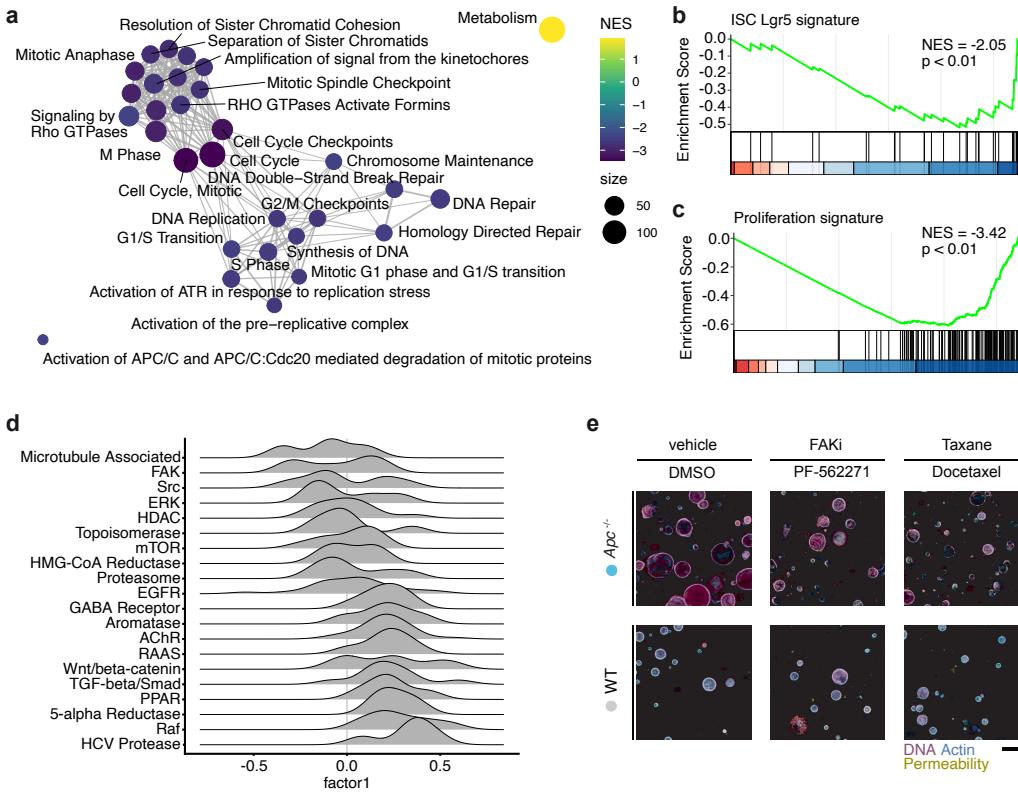


Figure 5.12: Factor 1, canonical Wnt signaling. a Gene-set enrichment network of factor 1 gene expression loadings. An edge connects Reactome pathways with more than 20% overlap. Central enriched processes include mitosis, DNA replication and DNA damage repair. b and c Gene set enrichment results of the "Lgr5 intestinal stem cell" and "proliferation" signature by Merlos-Suarez et al.¹⁸. over ranked factor 1 gene expression loadings (ranking from high factor 1 loading to low factor 1 loading, NES = normalized enrichment score). d Distributions of treatment activity loadings grouped by drug target for factor 1. e Example images of compound treated organoids with WT or Apc^{-/-} genotype. Representative images are displayed (magenta = DNA, cyan = actin, yellow = cell permeability, scale-bar: 200μm).

scores. Further exploration of the association between the treatment activity score and Apc genotype showed that small molecule inhibitors of the canonical Wnt secretion pathway protein Porcupine (Porcn), IWP-L6 and LGK-974, were more active in Apc WT organoids relative to their Apc^{-/-} counterparts¹⁵ (Figure 5.13a). In contrast, this effect was not observable for PRI-724, a small molecule inhibitor targeting the interaction of beta-catenin and CREB-binding-protein in the canonical Wnt signaling pathway²⁰ (Figure 5.13a).

The observed differences in treatment activity scores among small molecule inhibitors are most likely related to their different targets' relative location to Apc in the canonical Wnt signaling cascade. While

Porcn-dependent Wnt secretion is generally upstream of the destruction complex, the interaction of beta-catenin and the transcriptional coactivator CREB-binding-protein is located downstream of it. As a consequence, inhibition of destruction complex function by loss of Apc is expected to render cells less sensitive to perturbations of the Wnt secretion cascade than perturbations of transcription factor binding properties (Figure 5.13b).

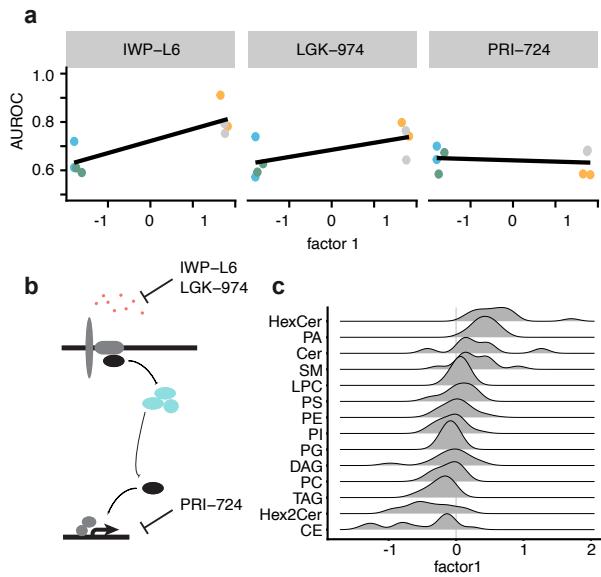


Figure 5.13: Small molecule Wnt signaling inhibitors. **a** AUROC activity score for three small molecule inhibitors of canonical Wnt signaling. **b** Target proteins for small molecules within the canonical Wnt signaling cascade with their relative position to the destruction complex (highlighted in blue). **c** Distributions of lipid abundance loadings grouped by lipid species for factor 1 (Abbreviations: HexCer - Hexosylceramide; PA - Phosphatidate; Cer - Ceramide; SM - Sphingomyelin; LPC - Lysophosphatidylcholine; PS - Phosphatidylserine; PE - Phosphatidylethanolamine; PI - Phosphatidylinositol; PG - Phosphatidylglycerol; DAG - Diacylglycerol; PC - Phosphatidylcholine; TAG - Triacylglycerol; Hex2Cer - Dihexosylceramide; CE - Cholesterolester)

Next to factor related differences in morphological compound sensitivity, I analyzed the association of factor 1 weights with measured lipid species. High concentrations of cholesterol esters were associated with a low factor 1 score (seen in Apc^{-/-} organoids), while elevated concentrations of phosphatidic acid species were linked to a high factor 1 score (figure 5.13c).

After linking factor 1 to Apc loss and identifying features caused by this molecular change, I was interested in identifying small molecule treatments that -based on the morphology they induced- shifted organoid state

along the factor 1 axis. As described in the previous chapter, small molecules that led to a predicted factor 1 change were identified using ANOVA. To identify treatments that led to a drop in organoid viability, a classifier developed by Jan Sauer and described in the previous chapter (LDC) trained on ground-truth lethal treatments within the stem-cell library was applied. Two groups of compounds were identified that induced a shift towards lower factor 1 scores (group A, figure 5.14) and higher factor 1 scores (group B).

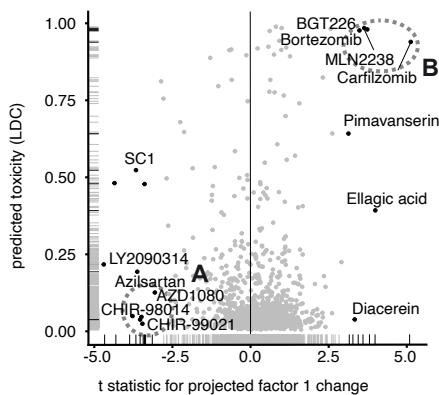


Figure 5.14: Projection of factor 1 scores for treatment-induced phenotypes and viability changes. Highlighted are compounds leading to a significant change in projected factor scores across all organoid lines (ANOVA). Organoid viability is predicted using a random-forest based classifier (LDC) with scores from 0 (no toxicity) to 1 (complete toxicity)

Small molecules within group A induced a morphology associated with Apc loss while maintaining organoid viability. In contrast, members of group B primarily led to a loss of viability and a shift towards a morphological state associated with Apc wildtype organoids. Of note, small molecules within these groups had related target proteins. Compounds within group A (incl. CHIR-98014, CHIR-99021, LY2090314) targeted GSK3 beta - a kinase with central function within the canonical Wnt signaling destruction complex. Inhibition of GSK3 beta leads to hyperactivation of canonical Wnt signaling. Members of group B primarily targeted the Proteasome (incl. Bortezomib, Carfilzomib, MLN2238).

Further validation of group A showed how treatment with the GSK3 beta inhibitor CHIR-98014 led to

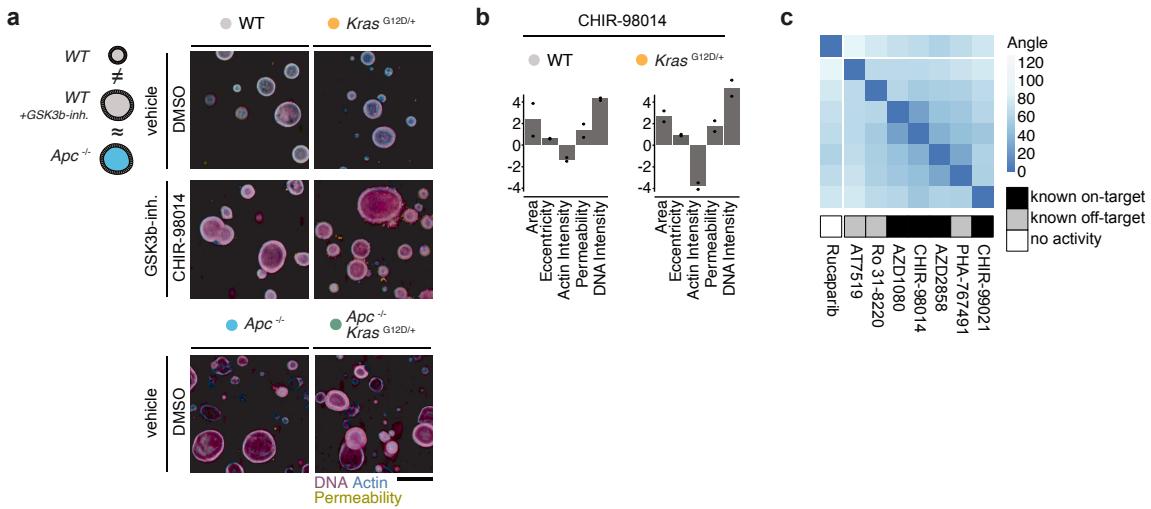


Figure 5.15: GSK3 beta inhibition dependent morphology in colon organoid models. **a** Small molecule inhibition of GSK3 beta (CHIR98014) leads to phenocopying of *Apc*^{-/-} genotype organoid models. **b** Shift of morphological features of wildtype and *Kras*^{G12D/+} organoid models treated with CHIR98014. Shown is an increase in organoid size (Area) and DNA intensity. **c** Excerpt of clustering from figure 5.9 d, labeled with known binding activity of listed small molecules. Rucaparib is not member of the cluster and shown for comparison.

treatment-induced phenotypes in WT and *Kras*^{G12D/+} organoids that phenocopied the unperturbed morphology of *Apc*^{-/-} and double mutant organoid models (Figure 5.15 a). On the feature level, treatment led to an increase in organoid size and DNA intensity (Figure 5.15 b) in *Apc* wildtype models. This change in morphology was likely due to an increased proliferation rate of mutant cells, leading to rising organoid size and a higher density of nuclei per analyzed object. Guided by the identification of a strong GSK3 beta inhibition phenotype in *Apc* wildtype organoids, I analyzed small molecules that clustered with known inhibitors of this kinase based on the angular distance of their drug effect vectors within these models (Figure 5.15 c) and 5.9 d). All small molecules clustering with well-described inhibitors of GSK3 beta had previously described off-target binding activity against this kinase within the LINCS KINOMEScan database⁶.

In conclusion of this section, loss of *Apc* function is the primary source of variation identified across the four analyzed organoid genotypes. Increased cell proliferation rate, canonical Wnt signaling and genomic stress caused by a loss of *Apc* is observable across profiling modalities and leads to characteristic dependencies (microtubuli, FAK signaling). Small molecules inhibiting the function of the destruction complex member GSK3 beta can phenocopy loss of *Apc* among colon organoid models.

5.8 AN ONCOGENE-INDUCED SENESCENCE PROGRAM CAUSED BY ISOLATED KrasG12D ACTIVATION

While the Apc^{-/-} genotype contributed primarily to factor 1 (Figure 5.14 b), the KrasG12D allele showed a strong loading for both factor 2 and factor 3. This observation paired with the fact that only organoid models without Apc loss of function were separated by factor 2 (Figure 5.10 c), led me to conclude that factor 2 described a KrasG12D dependent change in cell state in the presence of intact Apc function.

To understand the molecular mechanisms represented by factor 2, I again analyzed features with large absolute loadings for this factor. Plotting of factor loadings for treatment activity by small molecule target, showed that ERK and MEK inhibitors were more active in factor 2 low models (KrasG12D^{+/−}) while EGFR/HER2 inhibitors were more active in factor 2 high organoids (WT, figure 5.16 a and b). This juxtaposition in treatment activity against RAS-MAPK pathway members was reminiscent of the previous observations made around canonical Wnt signaling inhibitors (Figure 5.13). With oncogenic Kras localized between the receptor-layer (including Egfr and Her2) and downstream kinases (for example Erk), hyperactive Kras signaling likely leads to a cell state with relative resistance to EGFR inhibitors and increased dependency on Erk signaling. The previously observed transcriptional process of Egfr-downregulation as a response to KrasG12D^{+/−} is in line with these observations (Figure 5.5b).

In addition, HMG-CoA reductase inhibitors (incl. Rosuvastatin) showed an increased activity in KrasG12D^{+/−} organoids (Figure 5.16 a and b). This vulnerability towards cholesterol biosynthesis inhibitors in cells with oncogenic Kras signaling has previously been discussed³⁴.

Oncogene induced senescence is a cell state marked by an arrest of the cell cycle and expression of pro-inflammatory mediators as a response to an oncogenic perturbation. An activated KrasG12D^{+/−} genotype leads to oncogene induced senescence of colon epithelial cells *in vivo*².

Prompted by previous reports on the effect of oncogenic Kras, I identified an enrichment of a senescence related gene expression signature by Fridman et al. within the loadings of factor 2⁷ (Figure 5.16 c). Transcripts linked to cell senescence, including Igfbp3 (factor 2 loading ca. -0.999) and Hmgaz2 (factor 2 loading ca. -1.050) ranked among the strongest contributors to the factor.

Further exploration of loadings for factor 2 revealed systematic shifts in the composition of the lipidome.

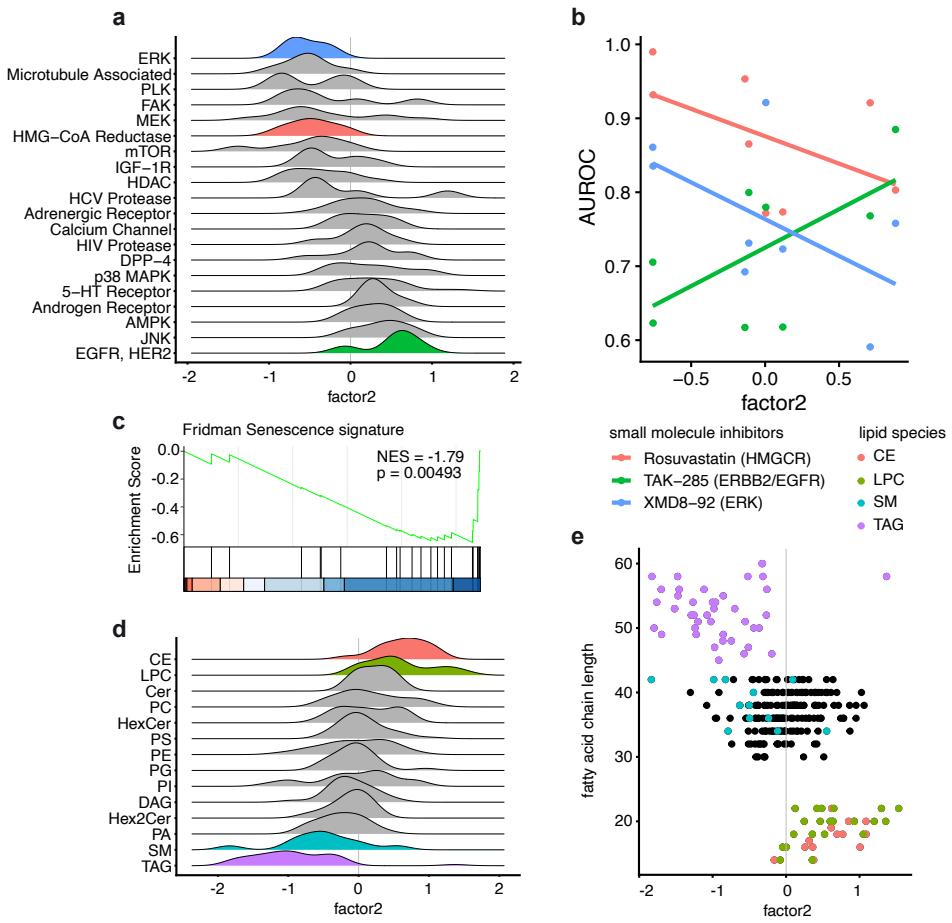


Figure 5.16: Factor 2, KrasG12D induced senescence. **a** Distributions of treatment activity loadings grouped by drug target for factor 2. **b** Relationship of representative drugs' activity with factor 2 score. Shown are compounds from highlighted groups in panel (a). **c** Gene set enrichment results of a senescence signature by ian et al. over ranked factor 2 gene expression loadings (ranking from high factor 2 loading to low factor 2 loading, NES = normalized enrichment score). **d** Distributions of lipid abundance loadings grouped by lipid species for factor 2 (Abbreviations: Hex-Cer - Hexosylceramide; PA - Phosphatidate; Cer - Ceramide; SM - Sphingomyelin; LPC - Lysophosphatidylcholine; PS - Phosphatidylserine; PE - Phosphatidylethanolamine; PI - Phosphatidylinositol; PG - Phosphatidylglycerol; DAG - Diacylglycerol; PC - Phosphatidylcholine; TAG - Triacylglycerol; Hex2Cer - Dihexosylceramide; CE - Cholesterolester). **e** Relationship of acyl chain length with factor 2 loading. Shown are lipids from highlighted species in panel (d)

Activation of oncogenic Kras led to an accumulation of sphingomyelin (SM) and triacylglycerol (TAG) species in colon cells. In contrast, I observed a relative depletion of cholesterol esters (CE) and lysophosphatidylcholine (LPC) (Figure 5.16 d). Next to shifts in the lipid composition, I also observed a correlation of fatty acid residue chain length and factor 2 loadings, where a low factor 2 score was correlated with higher residue chain length (Figure 5.16 e) and a higher degree of lipid desaturation (data not shown).

Both of these association was primarily driven by differences in the abundance of lipid species, however. To summarize, factor 2 explains differences between wildtype and KrasG12D+/- organoid models and captures the activation of RAS-MAPK signaling and subsequent oncogene induced senescence. KrasG12D+/- transformed colon cells showed a hyperactivation of RAS-MAPK signaling with characteristic changes in small molecule dependencies (ERK and EGFR inhibitors, respectively). Of note, KrasG12D+/- induced senescence coincided with the accumulation of triacylglycerol (TAG) and sphingomyelin (SM) species within colon organoids. Lipid accumulation in senescent cells has been observed in multiple senescent cells, including human fibroblasts (TAG accumulation)¹⁶ and, more specifically, has recently been described in non-mammalian model systems as a consequence of Kras mediated oncogene induced senescence³².

5.9 ONCOGENIC KRAS LEADS TO INCREASED mTOR SIGNALING IN THE CONTEXT OF APC LOSS OF FUNCTION

While factor 2 captured the effect of oncogenic Kras in an Apc wildtype state, factor 3 scores separated organoid models by Kras genotype in the context of an Apc loss of function (Figure 5.10 c). On average, factor 3 only accounted for ca. 10% of variance across modalities within the MOFA model, supporting the overall similarity of Apc^{-/-} and Apc^{-/-} KrasG12D^{+/+} organoids previously observed on the transcriptome, proteome, lipidome and phenotype level (Figure 5.4a-d and 5.7 b).

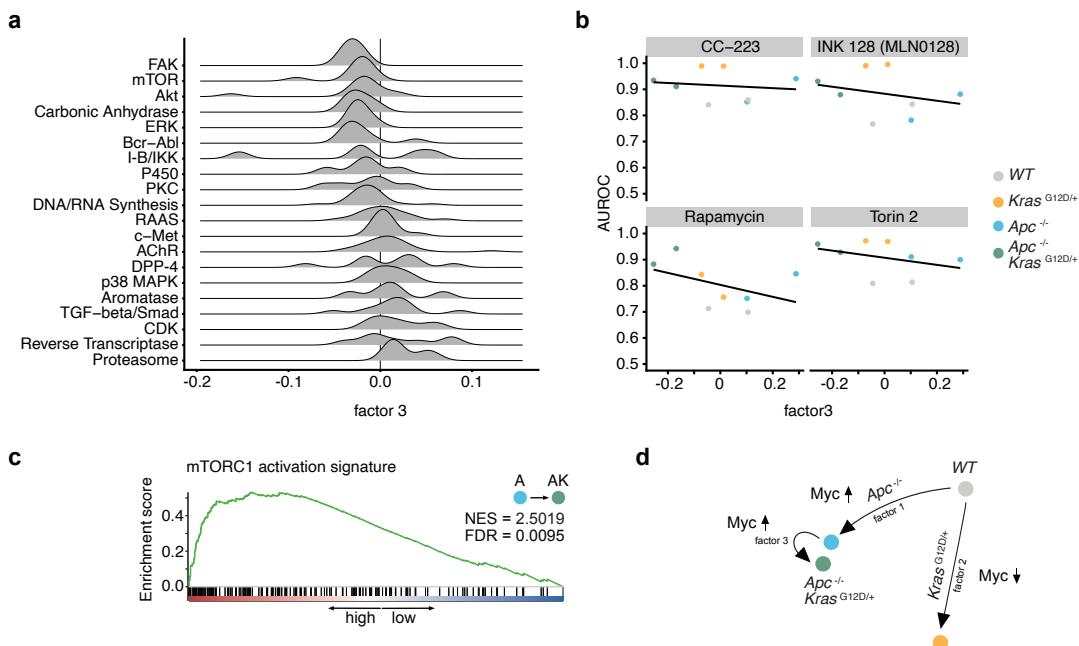


Figure 5.17: Factor 3, KrasG12D effects in the context of Apc loss of function. a Distributions of treatment activity loadings grouped by drug target for factor 3. b Relationship of representative mTOR inhibitor activity with factor 3 score. c Gene set enrichment results of a Reactome mTORC1 activation signature over ranked factor 3 gene expression loadings (ranking from high factor 3 loading to low factor 3 loading, NES = normalized enrichment score). d Visual summary of Myc gene set enrichment results for organoid state transitions. Myc signatures were significantly enriched in Apc^{-/-} models and depleted in models with isolated KrasG12D^{+/} mutation.

Exploration of factor 3 loadings identified mTOR and FAK inhibitors to systematically contribute to a negative factor score. Double-mutant organoid models (Apc^{-/-} KrasG12D^{+/}) showed a greater treatment activity score for these small molecule inhibitors than their single-mutant Apc^{-/-} counterparts (Figure 5.17 a).

This difference in treatment activity was observable for both ATP-competitive (e.g. INK-128, Sapanisertib) and non-ATP-competitive (e.g. Rapamycin) inhibitors (Figure 5.17 b). In line with the increased sensitivity to mTOR inhibitors, differential gene expression analysis comparing single (*Apc* $-/-$, abbreviated A) and double mutant (*Apc* $-/-$ *KrasG12D* $+/-$, abbreviated AK) organoids identified a significant increase in mTORC1 activation according to a Reactome signature (Figure 5.17 c, FDR=0.0095, NES=2.5019).

Further exploration of factor 3 loadings in gene expression space showed an enrichment of multiple Myc target gene signatures^{33,35}. Transcription signatures associated with increased Myc activity are enriched in double-mutant organoid models (*Apc* $-/-$ *KrasG12D* $+/-$) when compared to single-mutant counterparts (Figure 5.17 d). In fact, further analysis of factor 1 and factor 2 gene expression loadings identified significant changes in Myc activity signatures for both factors as well. Simplified, loss of function of *Apc* led to an increased activity of Myc (factor 1) while the isolated presence of oncogenic *KrasG12D* $+/-$ led to a reduced activity of Myc. These observations are consistent with previous studies describing a central role of increased Myc activity in Wnt-signaling dependent adenoma formation²⁶ and decreased Myc activity in oncogene induced senescence³².

In summary, factor 3 is representing the interaction of *Apc* loss of function and oncogenic *KrasG12D* $+/-$. In the mouse colon organoid model, the introduction of oncogenic *KrasG12D* $+/-$ within an *Apc* $-/-$ background leads to an increased mTORC1 signaling related gene expression signature and increased sensitivity to small molecule mTOR inhibitors. Myc activity related gene expression is increased in all factors related to malignant transformation, except the introduction of isolated oncogenic *KrasG12D* $+/-$ in where Myc activity is reduced in line with the oncogene induced senescence model.

5.10 ELLAGIC ACID IS A CANDIDATE SMALL MOLECULE REVERTANT THERAPEUTIC OF ORGANOID ADENOMA STATES

Similar to the analysis in the previous chapter, I searched for treatments which induced organoid phenotypes that suggested a change of organoid state within the learned factor-space. Treatments that were predicted to move organoids towards a wildtype state along factor 1 and 2 were of particular medical relevance. Analogous to the concept of revertant genetic mutations, I referred to small molecules with a predicted ability to move transformed organoids towards a wildtype state as potential revertant therapeutics.

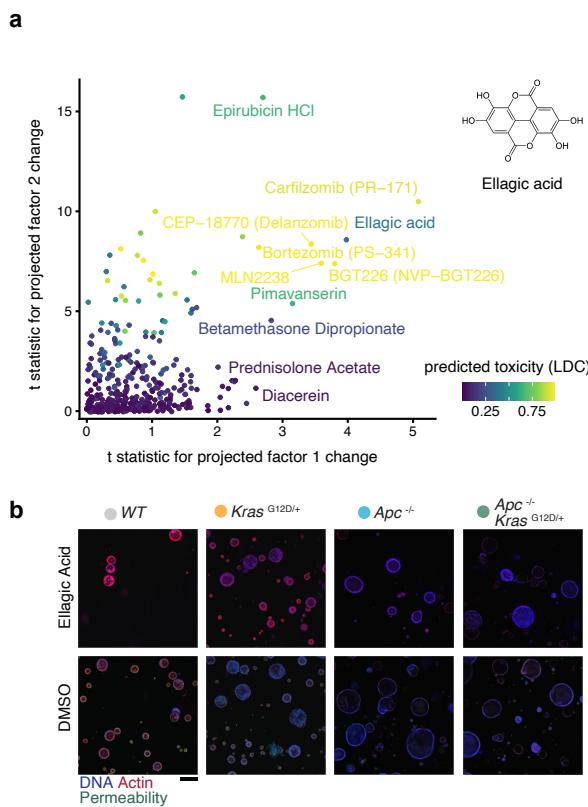


Figure 5.18: Projected changes in factor 1 and 2 scores after small molecule treatment **a** Shows are treatments leading to change in projected factor scores across all organoid lines (ANOVA). Color coding represents predictions based on LDC viability classifier. The structural formula of Ellagic Acid is depicted in the top right. **b** Organoid treatment with Ellagic acid. Depicted are representative example images for each line (blue = DNA, red = actin, green = cell permeability, scale bar=200μm).

The majority of compounds that led to a predicted shift towards a wildtype organoid state along both axes were also associated with a high predicted toxicity (Figure 5.18 a, color coding are predictions based on LDC viability classifier). Ellagic acid, however, a natural compound that was predicted to function as a revertant, showed a moderate predicted toxicity (Figure 5.18 a, top right). Inspection of organoid models treated with Ellagic acid showed reduced number and size of organoids compared to the vehicle control, while organoid viability was generally maintained (Figure 5.18 b).

In this chapter I hypothesized that image-based profiling of engineered organoid models can help identify small molecule candidates that move organoids within a well-defined representation space. Analysis of four genetically defined organoid models resulted in three interpretable factors within the MOFA analysis.

Exploration of factors revealed that Kras G12D dependent oncogene induced senescence in colon organoids is accompanied with characteristic changes to the lipidome, which are not observable in Apc mutant organoid models.

The proposed approach led to the identification of both a well-described and potentially novel set of small molecules that can move organoids in the learned representation space. Known small molecule inhibitors of GSK₃ beta were predicted to move organoid in factor space along factor 1, a canonical Wnt-signaling associated program. Ellagic acid, a natural compound, was predicted to move organoid phenotypes towards an unperturbed morphology, prompting further validation.

References

- [noa] MaxContrastProjection. <https://bioconductor.riken.jp/packages/3.10/bioc/html/MaxContrastProjection.html>. Accessed: 2022-8-28.
- [2] Bennecke, M., Kriegl, L., Bajbouj, M., Retzlaff, K., Robine, S., Jung, A., Arkan, M. C., Kirchner, T., & Greten, F. R. (2010). Ink4a/Arf and oncogene-induced senescence prevent tumor progression during alternative colorectal tumorigenesis. *Cancer cell*, 18(2), 135–146.
- [3] Betge, J., Rindtorff, N., Sauer, J., Rauscher, B., Dingert, C., Gaitanzi, H., Herweck, F., Srour-Mhanna, K., Miersch, T., Valentini, E., Boonekamp, K. E., Hauber, V., Gutting, T., Frank, L., Belle, S., Gaiser, T., Buchholz, I., Jesenofsky, R., Härtel, N., Zhan, T., Fischer, B., Breitkopf-Heinlein, K., Burgermeister, E., Ebert, M. P., & Boutros, M. (2022). The drug-induced phenotypic landscape of colorectal cancer organoids. *Nature communications*, 13(1), 3135.
- [4] Broutier, L., Mastrogiovanni, G., Verstegen, M. M. A., Francies, H. E., Gavarró, L. M., Bradshaw, C. R., Allen, G. E., Arnes-Benito, R., Sidorova, O., Gaspersz, M. P., Georgakopoulos, N., Koo, B.-K., Dietmann, S., Davies, S. E., Praseedom, R. K., Lieshout, R., IJzermans, J. N. M., Wigmore, S. J., Saeb-Parsy, K., Garnett, M. J., van der Laan, L. J. W., & Huch, M. (2017). Human primary liver cancer-derived organoid cultures for disease modeling and drug screening. *Nature medicine*, 23(12), 1424–1435.
- [5] Drost, J., van Jaarsveld, R. H., Ponsioen, B., Zimberlin, C., van Boxtel, R., Buijs, A., Sachs, N., Overmeer, R. M., Offerhaus, G. J., Begthel, H., Korving, J., van de Wetering, M., Schwank, G., Logtenberg, M., Cuppen, E., Snippert, H. J., Medema, J. P., Kops, G. J. P. L., & Clevers, H. (2015). Sequential cancer mutations in cultured human intestinal stem cells. *Nature*, 521(7550), 43–47.
- [6] Duan, Q., Flynn, C., Niepel, M., Hafner, M., Muhlich, J. L., Fernandez, N. F., Rouillard, A. D., Tan, C. M., Chen, E. Y., Golub, T. R., Sorger, P. K., Subramanian, A., & Ma'ayan, A. (2014). LINCS canvas browser: interactive web app to query, browse and interrogate LINCS L1000 gene expression signatures. *Nucleic acids research*, 42(Web Server issue), W449–60.
- [7] Fridman, A. L. & Tainsky, M. A. (2008). Critical pathways in cellular senescence and immortalization revealed by gene expression profiling. *Oncogene*, 27(46), 5975–5987.
- [8] Fujii, M., Shimokawa, M., Date, S., Takano, A., Matano, M., Nanki, K., Ohta, Y., Toshimitsu, K., Nakazato, Y., Kawasaki, K., Uraoka, T., Watanabe, T., Kanai, T., & Sato, T. (2016). A colorectal tumor organoid library demonstrates progressive loss of niche factor requirements during tumorigenesis. *Cell stem cell*, 18(6), 827–838.
- [9] Griss, J., Viteri, G., Sidiroopoulos, K., Nguyen, V., Fabregat, A., & Hermjakob, H. (2020). ReactomeGSA - efficient Multi-Omics comparative pathway analysis. *Molecular & cellular proteomics: MCP*, 19(12), 2115–2125.

- [10] Guinney, J., Dienstmann, R., Wang, X., de Reyniès, A., Schlicker, A., Soneson, C., Marisa, L., Roepman, P., Nyamundanda, G., Angelino, P., Bot, B. M., Morris, J. S., Simon, I. M., Gerster, S., Fessler, E., De Sousa E Melo, F., Missaglia, E., Ramay, H., Barras, D., Homicsko, K., Maru, D., Manyam, G. C., Broom, B., Boige, V., Perez-Villamil, B., Laderas, T., Salazar, R., Gray, J. W., Hanahan, D., Tabernero, J., Bernards, R., Friend, S. H., Laurent-Puig, P., Medema, J. P., Sadanandam, A., Wessels, L., Delorenzi, M., Kopetz, S., Vermeulen, L., & Tejpar, S. (2015). The consensus molecular subtypes of colorectal cancer. *Nature medicine*, 21(11), 1350–1356.
- [11] Isella, C., Brundu, F., Bellomo, S. E., Galimi, F., Zanella, E., Porporato, R., Petti, C., Fiori, A., Orzan, F., Senetta, R., Boccaccio, C., Ficarra, E., Marchionni, L., Trusolino, L., Medico, E., & Bertotti, A. (2017). Selective analysis of cancer-cell intrinsic transcriptional traits defines novel clinically relevant subtypes of colorectal cancer. *Nature communications*, 8(1), 15107.
- [12] Jackson, E. L., Willis, N., Mercer, K., Bronson, R. T., Crowley, D., Montoya, R., Jacks, T., & Tuvesson, D. A. (2001). Analysis of lung tumor initiation and progression using conditional expression of oncogenic k-ras. *Genes & development*, 15(24), 3243–3248.
- [13] Klaeger, S., Heinzelmeir, S., Wilhelm, M., Polzer, H., Vick, B., Koenig, P.-A., Reinecke, M., Ruprecht, B., Petzoldt, S., Meng, C., Zecha, J., Reiter, K., Qiao, H., Helm, D., Koch, H., Schoof, M., Canevari, G., Casale, E., Depaolini, S. R., Feuchtinger, A., Wu, Z., Schmidt, T., Rueckert, L., Becker, W., Huenges, J., Garz, A.-K., Gohlke, B.-O., Zolg, D. P., Kayser, G., Voorder, T., Preissner, R., Hahne, H., Tönisson, N., Kramer, K., Götzke, K., Bassermann, F., Schlegl, J., Ehrlich, H.-C., Aiche, S., Walch, A., Greif, P. A., Schneider, S., Felder, E. R., Ruland, J., Médard, G., Jeremias, I., Spiekermann, K., & Kuster, B. (2017). The target landscape of clinical kinase drugs. *Science*, 358(6367).
- [14] Kotliarova, S., Pastorino, S., Kovell, L. C., Kotliarov, Y., Song, H., Zhang, W., Bailey, R., Maric, D., Zenklusen, J. C., Lee, J., & Fine, H. A. (2008). Glycogen synthase kinase-3 inhibition induces glioma cell death through c-MYC, nuclear factor-kappab, and glucose regulation. *Cancer research*, 68(16), 6642–6651.
- [15] Liu, J., Pan, S., Hsieh, M. H., Ng, N., Sun, F., Wang, T., Kasibhatla, S., Schuller, A. G., Li, A. G., Cheng, D., Li, J., Tompkins, C., Pferdekamper, A., Steffy, A., Cheng, J., Kowal, C., Phung, V., Guo, G., Wang, Y., Graham, M. P., Flynn, S., Brenner, J. C., Li, C., Villarroel, M. C., Schultz, P. G., Wu, X., McNamara, P., Sellers, W. R., Petruzzelli, L., Boral, A. L., Seidel, H. M., McLaughlin, M. E., Che, J., Carey, T. E., Vanasse, G., & Harris, J. L. (2013). Targeting wnt-driven cancer through the inhibition of porcupine by LGK974. *Proceedings of the National Academy of Sciences of the United States of America*, 110(50), 20224–20229.
- [16] Lizardo, D. Y., Lin, Y.-L., Gokcumen, O., & Atilla-Gokcumen, G. E. (2017). Regulation of lipids is central to replicative senescence. *Molecular bioSystems*, 13(3), 498–509.
- [17] Matano, M., Date, S., Shimokawa, M., Takano, A., Fujii, M., Ohta, Y., Watanabe, T., Kanai, T., & Sato, T. (2015). Modeling colorectal cancer using CRISPR-Cas9-mediated engineering of human intestinal organoids. *Nature medicine*, 21(3), 256–262.
- [18] Merlos-Suárez, A., Barriga, F. M., Jung, P., Iglesias, M., Céspedes, M. V., Rossell, D., Sevillano, M., Hernando-Momblona, X., da Silva-Diz, V., Muñoz, P., Clevers, H., Sancho, E., Mangues, R., & Batlle, E. (2011). The intestinal stem cell signature identifies colorectal cancer stem cells and predicts disease relapse. *Cell stem cell*, 8(5), 511–524.

- [19] Muzny, D. M., Bainbridge, M. N., Chang, K., Dinh, H. H., Drummond, J. A., Fowler, G., Kovar, C. L., Lewis, L. R., Morgan, M. B., Newsham, I. F., Reid, J. G., Santibanez, J., Shinbrot, E., Trevino, L. R., Wu, Y.-Q., Wang, M., Gunaratne, P., Donehower, L. A., Creighton, C. J., Wheeler, D. A., Gibbs, R. A., Lawrence, M. S., Voet, D., Jing, R., Cibulskis, K., Sivachenko, A., Stojanov, P., McKenna, A., Lander, E. S., Gabriel, S., Getz, G., Ding, L., Fulton, R. S., Koboldt, D. C., Wylie, T., Walker, J., Dooling, D. J., Fulton, L., Delehaunty, K. D., Fronick, C. C., Demeter, R., Mardis, E. R., Wilson, R. K., Chu, A., Chun, H.-J. E., Mungall, A. J., Pleasance, E., Gordon Robertson, A., Stoll, D., Balasundaram, M., Birol, I., Butterfield, Y. S. N., Chuah, E., Cope, R. J. N., Dhalla, N., Guin, R., Hirst, C., Hirst, M., Holt, R. A., Lee, D., Li, H. I., Mayo, M., Moore, R. A., Schein, J. E., Slobodan, J. R., Tam, A., Thiessen, N., Varhol, R., Zeng, T., Zhao, Y., Jones, S. J. M., Marra, M. A., Bass, A. J., Ramos, A. H., Saksena, G., Cherniack, A. D., Schumacher, S. E., Tabak, B., Carter, S. L., Pho, N. H., Nguyen, H., Onofrio, R. C., Crenshaw, A., Ardlie, K., Beroukhim, R., Winckler, W., Getz, G., Meyerson, M., Protopopov, A., Zhang, J., Hadjipanayis, A., Lee, E., Xi, R., Yang, L., Ren, X., Zhang, H., Sathiamoorthy, N., Shukla, S., Chen, P.-C., Haseley, P., Xiao, Y., Lee, S., Seidman, J., Chin, L., Park, P. J., Kucherlapati, R., Todd Auman, J., Hoadley, K. A., Du, Y., Wilkerson, M. D., Shi, Y., Liquori, C., Meng, S., Li, L., Turman, Y. J., Topal, M. D., Tan, D., Waring, S., Buda, E., Walsh, J., Jones, C. D., Mieczkowski, P. A., Singh, D., Wu, J., Gulabani, A., Dolina, P., Bodenheimer, T., Hoyle, A. P., Simons, J. V., Soloway, M., Mose, L. E., Jefferys, S. R., Balu, S., O'Connor, B. D., Prins, J. F., Chiang, D. Y., Neil Hayes, D., Perou, C. M., Hinoue, T., Weisenberger, D. J., Maglinte, D. T., Pan, F., Berman, B. P., Van Den Berg, D. J., Shen, H., Triche, Jr, T., Baylin, S. B., Laird, P. W., Getz, G., Noble, M., Voet, D., Saksena, G., Gehlenborg, N., DiCaro, D., Zhang, J., Zhang, H., Wu, C.-J., Yingchun Liu, S., Shukla, S., Lawrence, M. S., Zhou, L., Sivachenko, A., Lin, P., Stojanov, P., Jing, R., Park, R. W., Nazaire, M.-D., Robinson, J., Thorvaldsdottir, H., Mesirov, J., Park, P. J., Chin, L., Thorsson, V., Reynolds, S. M., Bernard, B., Kreisberg, R., Lin, J., Iype, L., Bressler, R., Erkkilä, T., Gundapuneni, M., Liu, Y., Norberg, A., Robinson, T., Yang, D., Zhang, W., Shmulevich, I., de Ronde, J. J., Schultz, N., Cerami, E., Ciriello, G., Goldberg, A. P., Gross, B., Jacobsen, A., Gao, J., Kaczkowski, B., Sinha, R., Arman Aksoy, B., Antipin, Y., Reva, B., Shen, R., Taylor, B. S., Chan, T. A., Ladanyi, M., Sander, C., Akbani, R., Zhang, N., Broom, B. M., Casasent, T., Unruh, A., Wakefield, C., Hamilton, S. R., Craig Cason, R., Baggerly, K. A., Weinstein, J. N., Haussler, D., Benz, C. C., Stuart, J. M., Benz, S. C., Zachary Sanborn, J., Vaske, C. J., Zhu, J., Szeto, C., Scott, G. K., Yau, C., Ng, S., Goldstein, T., Ellrott, K., Collisson, E., Cozen, A. E., Zerbino, D., Wilks, C., Craft, B., Spellman, P., Penny, R., Shelton, T., Hatfield, M., Morris, S., Yena, P., Shelton, C., Sherman, M., Paulauskis, J., Gastier-Foster, J. M., Bowen, J., Ramirez, N. C., Black, A., Pyatt, R., Wise, L., White, P., Bertagnolli, M., Brown, J., Chan, T. A., Chu, G. C., Czerwinski, C., Denstman, F., Dhir, R., Dörner, A., Fuchs, C. S., Guillem, J. G., Iaccoca, M., Juhl, H., Kaufman, A., Kohl, III, B., Van Le, X., Mariano, M. C., Medina, E. N., Meyers, M., Nash, G. M., Paty, P. B., Petrelli, N., Rabeno, B., Richards, W. G., Solit, D., Swanson, P., Temple, L., Tepper, J. E., Thorp, R., Vakiani, E., Weiser, M. R., Willis, J. E., Witkin, G., Zeng, Z., Zinner, M. J., Zornig, C., Jensen, M. A., Sfeir, R., Kahn, A. B., Chu, A. L., Kothiyal, P., Wang, Z., Snyder, E. E., Pontius, J., Pihl, T. D., Ayala, B., Backus, M., Walton, J., Whitmore, J., Baboud, J., Bertron, D. L., Nicholls, M. C., Srinivasan, D., Raman, R., Girshik, S., Kigonya, P. A., Alonso, S., Sanbhadt, R. N., Barletta, S. P., Greene, J. M., Pot, D. A., Mills Shaw, K. R., Dillon, L. A. L., Buetow, K., Davidsen, T., Demchok, J. A., Eley, G., Ferguson, M., Fielding, P., Schaefer, C., Sheth, M., Yang, L., Guyer, M. S., Ozenberger, B. A., Palchik, J. D., Peterson, J., Sofia, H. J., & Thomson, E. (2012). Comprehensive molecular characterization of human colon and rectal cancer. *Nature*, 487(7407), 330–337.
- [20] Okazaki, H., Sato, S., Koyama, K., Morizumi, S., Abe, S., Azuma, M., Chen, Y., Goto, H., Aono, Y., Ogawa, H., Kagawa, K., Nishimura, H., Kawano, H., Toyoda, Y., Uehara, H., Kouji, H., & Nishioka,

- Y. (2019). The novel inhibitor PRI-724 for Wnt/β-catenin/CBP signaling ameliorates bleomycin-induced pulmonary fibrosis in mice. *Experimental lung research*, 45(7), 188–199.
- [21] O'Reilly, K. E., Rojo, F., She, Q.-B., Solit, D., Mills, G. B., Smith, D., Lane, H., Hofmann, F., Hicklin, D. J., Ludwig, D. L., Baselga, J., & Rosen, N. (2006). mTOR inhibition induces upstream receptor tyrosine kinase signaling and activates akt. *Cancer research*, 66(3), 1500–1508.
- [22] Patankar, M., Eskelinan, S., Tuomisto, A., Mäkinen, M. J., & Karttunen, T. J. (2019). KRAS and BRAF mutations induce anoikis resistance and characteristic 3D phenotypes in caco-2 cells. *Molecular medicine reports*, 20(5), 4634–4644.
- [23] Sansom, O. J., Meniel, V. S., Muncan, V., Phesse, T. J., Wilkins, J. A., Reed, K. R., Vass, J. K., Athineos, D., Clevers, H., & Clarke, A. R. (2007). Myc deletion rescues apc deficiency in the small intestine. *Nature*, 446(7136), 676–679.
- [24] Sato, T., Stange, D. E., Ferrante, M., Vries, R. G. J., Van Es, J. H., Van den Brink, S., Van Houdt, W. J., Pronk, A., Van Gorp, J., Siersema, P. D., & Clevers, H. (2011). Long-term expansion of epithelial organoids from human colon, adenoma, adenocarcinoma, and barrett's epithelium. *Gastroenterology*, 141(5), 1762–1772.
- [25] Sato, T., Vries, R. G., Snippert, H. J., van de Wetering, M., Barker, N., Stange, D. E., van Es, J. H., Abo, A., Kujala, P., Peters, P. J., & Clevers, H. (2009). Single lgr5 stem cells build crypt-villus structures in vitro without a mesenchymal niche. *Nature*, 459(7244), 262–265.
- [26] Satoh, K., Yachida, S., Sugimoto, M., Oshima, M., Nakagawa, T., Akamoto, S., Tabata, S., Saitoh, K., Kato, K., Sato, S., Igarashi, K., Aizawa, Y., Kajino-Sakamoto, R., Kojima, Y., Fujishita, T., Enomoto, A., Hirayama, A., Ishikawa, T., Taketo, M. M., Kushida, Y., Haba, R., Okano, K., Tomita, M., Suzuki, Y., Fukuda, S., Aoki, M., & Soga, T. (2017). Global metabolic reprogramming of colorectal cancer occurs at adenoma stage and is induced by MYC. *Proceedings of the National Academy of Sciences of the United States of America*, 114(37), E7697–E7706.
- [27] Schell, M. J., Yang, M., Teer, J. K., Lo, F. Y., Madan, A., Coppola, D., Monteiro, A. N. A., Nebozhyn, M. V., Yue, B., Loboda, A., Bien-Willner, G. A., Greenawalt, D. M., & Yeatman, T. J. (2016). A multi-gene mutation classification of 468 colorectal cancers reveals a prognostic role for APC. *Nature communications*, 7, 11743.
- [28] Schütte, M., Risch, T., Abdavi-Azar, N., Boehnke, K., Schumacher, D., Keil, M., Yildirim, R., Jan-drasis, C., Borodina, T., Amstislavskiy, V., Worth, C. L., Schweiger, C., Liebs, S., Lange, M., Warnatz, H.-J., Butcher, L. M., Barrett, J. E., Sultan, M., Wierling, C., Golob-Schwarzl, N., Lax, S., Uranitsch, S., Becker, M., Welte, Y., Regan, J. L., Silvestrov, M., Kehler, I., Fusi, A., Kessler, T., Herwig, R., Landegren, U., Wienke, D., Nilsson, M., Velasco, J. A., Garin-Chesa, P., Reinhard, C., Beck, S., Schäfer, R., Regenbrecht, C. R. A., Henderson, D., Lange, B., Haybaeck, J., Keilholz, U., Hoffmann, J., Lehrach, H., & Yaspo, M.-L. (2017). Molecular dissection of colorectal cancer in pre-clinical models identifies biomarkers predicting sensitivity to EGFR inhibitors. *Nature communications*, 8, 14262.
- [29] Sharma, S. V., Lee, D. Y., Li, B., Quinlan, M. P., Takahashi, F., Maheswaran, S., McDermott, U., Azizian, N., Zou, L., Fischbach, M. A., Wong, K.-K., Brandstetter, K., Wittner, B., Ramaswamy, S., Clas-sion, M., & Settleman, J. (2010). A chromatin-mediated reversible drug-tolerant state in cancer cell subpopulations. *Cell*, 141(1), 69–80.

- [30] Van De Wetering, M., Francies, H. E., Francis, J. M., Bounova, G., Iorio, F., Pronk, A., Van Houdt, W., Van Gorp, J., Taylor-Weiner, A., Kester, L., McLaren-Douglas, A., Blokker, J., Jaksani, S., Bartfeld, S., Volckman, R., Van Sluis, P., Li, V. S. W., Seepo, S., Sekhar Pedamallu, C., Cibulskis, K., Carter, S. L., McKenna, A., Lawrence, M. S., Lichtenstein, L., Stewart, C., Koster, J., Versteeg, R., Van Oudenaarden, A., Saez-Rodriguez, J., Vries, R. G. J., Getz, G., Wessels, L., Stratton, M. R., McDermott, U., Meyerson, M., Garnett, M. J., & Clevers, H. (2015). Prospective derivation of a living organoid biobank of colorectal cancer patients. *Cell*, 161(4), 933–945.
- [31] Weeber, F., van de Wetering, M., Hoogstraat, M., Dijkstra, K. K., Krijgsman, O., Kuilman, T., Gadella-van Hooijdonk, C. G. M., van der Velden, D. L., Peper, D. S., Cuppen, E. P. J. G., Vries, R. G., Clevers, H., & Voest, E. E. (2015). Preserved genetic diversity in organoids cultured from biopsies of human colorectal cancer metastases. *Proceedings of the National Academy of Sciences*, 112(43), 201516689.
- [32] Yao, Y., Sun, S., Wang, J., Fei, F., Dong, Z., Ke, A.-W., He, R., Wang, L., Zhang, L., Ji, M.-B., Li, Q., Yu, M., Shi, G.-M., Fan, J., Gong, Z., & Wang, X. (2018). Canonical wnt signaling remodels lipid metabolism in zebrafish hepatocytes following ras oncogenic insult. *Cancer research*, 78(19), 5548–5560.
- [33] Yu, D., Cozma, D., Park, A., & Thomas-Tikhonenko, A. (2005). Functional validation of genes implicated in lymphomagenesis: an in vivo selection assay using a myc-induced b-cell tumor. *Annals of the New York Academy of Sciences*, 1059, 145–159.
- [34] Yu, R., Longo, J., van Leeuwen, J. E., Mullen, P. J., Ba-Alawi, W., Haibe-Kains, B., & Penn, L. Z. (2018). Statin-Induced cancer cell death can be mechanistically uncoupled from prenylation of RAS family proteins. *Cancer research*, 78(5), 1347–1357.
- [35] Zeller, K. I., Jegga, A. G., Aronow, B. J., O'Donnell, K. A., & Dang, C. V. (2003). An integrated database of genes responsive to the myc oncogenic transcription factor: identification of direct genomic targets. *Genome biology*, 4(10), R69.

EIGENANTEIL AN DATENERHEBUNG UND -AUSWERTUNG SOWIE EIGENE
VERÖFFENTLICHUNGEN

MANUSKRIPTE, WELCHE IN DIESER DISSERTATION BEHANDELT WERDEN:

1. BETGE J*, Rindtorff N*, SAUER J*, RAUSCHER B, DINGERT C, GAITANTZI H, HERWECK F, MIER SCH T, VALENTINI E, HAUBER V, GUTTING T, FRANK L, BELLE S, GAISER T, BUCHHOLZ I, JESENOFSKY R, HAERTEL N, ZHAN T, FISCHER B, BREITKOPF-HEINLEIN K, BURGERMEISTER E, EBERT MP AND BOUTROS M. AUTOMATED IMAGE-BASED PROFILING IDENTIFIES DISTINCT COMPOUND INDUCED PHENOTYPES IN PATIENT-DERIVED ORGANOID S. MANUSCRIPT IN PREPARATION.
2. Rindtorff N, SAUER J, BETGE J, ZHAN, T, BOHMANN N, FISCHER B., SCHOLL C, BOUTROS M.
MANUSCRIPT IN PREPARATION

IM RAHMEN DER ERSTGENANNTEN ARBEIT ETABLIERTE ICH DIE METHODIK ZUR KULTIVIERUNG UND HOCHDURCHSATZ-MEDIKAMENTENTESTUNG VON HUMANEN DARMKREBSORGANOÏDEN IN DER ARBEITSGRUPPE VON MICHAEL BOUTROS. VON BEGINN DES PROJEKTES AN WAR ICH MIT JOHANNES BETGE FÜR DIE ZIELSETZUNG UND PLANUNG VERANTWORTLICH. ZUSAMMEN MIT HERRN BETGE, INITIIERTE UND BEGLEITETE WIR DIE ANALYSE DER MESSERGEBNISSE DURCH JAN SAUER UND, KURZ VOR EINREICHEN DER ERSTEN FASSUNG DES MANUSKRIPTES, AUCH DURCH BENEDIKT RAUSCHER. INSGESAMT BELÄUFT SICH MEIN BEITRAG ZU DIESEM MANUSKRIFT AUF 30-40% DER GESAMTLEISTUNG.

IM RAHMEN DER ZWEITGENANNTEN ARBEIT FÜHRTE ICH DIE KONSEPTION, ETABLIERUNG, EXEKUTION UND DATENANALYSE DES PROJEKTES KOMPLETT SELBSTSTÄNDIG DURCH. FÜR TEILE DER BILDANALYSE NUTZE ICH VON JAN SAUER BEREITES ENTWICKELTE METHODEN. JOHANNES BETGE UND TIANZUO ZHAN DIENTEN ALS BERATER WÄHREND DER INITIALEN PHASE DES PROJEKTES. NICK BOHMANN ARBEITETE ALS EIN BESUCHENDER STUDENT FÜR 6 WOCHE N AUF DEM PROJEKT MIT. BERND FISCHER WAR VOR SEINEM TOD DER ARBEITSGRUPPENLEITER VON JAN SAUER. CLAUDIA SCHOLLS ARBEITSGRUPPE STELLTE DAS MAUSMODELL ZUR VERFÜGUNG. MICHAEL BOUTROS WAR BETREUENDER DOKTORVATER.

WEITERE KOLLABORATIONEN:

1. HEIGWER F, SCHEEDER C, LAUFER C, RAUSCHER B, Rindtorff N, MIERSCH T, SCHMITT B, BLASS C, BOUTROS M. A HIGH-RESOLUTION MAP OF SIGNALING ACROSS THE METAZOAN CELL. SUBMITTED.
2. ZHAN T*, AMBROSI G*, WANDMACHER MA*, RAUSCHER B, BETGE J, Rindtorff N, HAEUSSLER R, HINSENKAMP I, BAMBERG L, HESSLING B, ERDMANN G, BURGERMEISTER E, EBERT MP AND BOUTROS M. MEK INHIBITION ACTIVATES WNT SIGNALLING AND INDUCES STEM CELL PLASTICITY IN COLORECTAL CANCER ORGANOIDS. NATURE COMMUNICATIONS - 2018
3. ZHAN T*, Rindtorff N*, BETGE J, EBERT M, BOUTROS M. CRISPR/CAS9 FOR CANCER RESEARCH AND THERAPY. SEMINARS IN CANCER BIOLOGY - 2017
4. ZHAN T*, RINDTORFF N*, BOUTROS M. WNT SIGNALING IN CANCER. ONCOGENE. - 2016

* EQUAL CONTRIBUTION

Danksagung

ZUM ENDE DIESER DISSERTATION MÖCHTE ICH MICH BEI EINER REIHE VON MENSCHEN BEDANKEN, die mich während der letzten Jahre begleitet haben. Nicht alle Gelisteten sind inzwischen noch unter uns und zu manchen habe ich leider nur noch wenig Kontakt.

EIDESSTATTLICHE VERSICHERUNG

1. BEI DER EINGEREICHTEN DISSERTATION ZU DEM THEMA "MULTIPARAMETRIC PHENOTYPING OF INTESTINAL ORGANOIDS TO MODEL DISEASE INITIATION AND TREATMENT REONSE IN COLORECTAL CANCER" HANDELT ES SICH UM MEINE EIGENSTÄNDIG ERBRACHTE LEISTUNG.
2. ICH HABE NUR DIE ANGEgebenEN QUELLEN UND HILFSMITTEL BENUTZT UND MICH KEINER UNZULÄSSIGEN HILFE DRITTER BEDIENT. INSbesondere HABE ICH WÖRTLICH ODER SINNGEMÄSS AUS ANDEREN WERKEN ÜBERNOMMENE INHALTE ALS SOLCHE KENNTLICH GEMACHT.
3. DIE ARBEIT ODER TEILE DAVON HABE ICH BISLANG NICHT AN EINER HOCHSCHULE DES IN- ODER AUSLANDS ALS BESTANDTEIL EINER PRÜFUNGS- ODER QUALIFIKATIONSLEISTUNG VORGELEGT.
4. DIE RICHTIGKEIT DER VORSTEHENDEN ERKLÄRUNGEN BESTÄTIGE ICH.
5. DIE BEDEUTUNG DER EIDESSTATTLICHEN VERSICHERUNG UND DIE STRAFRECHTLICHEN FOLGEN EINER UNRICHTIGEN ODER UNVOLLSTÄNDIGEN EIDESSTATTLICHEN VERSICHERUNG SIND MIR BEKANNT. ICH VERSICHERE AN EIDES STATT, DASS ICH NACH BESTEM WISSEN DIE REINE WAHRHEIT ERKLÄRT UND NICHTS VERSCHWIEGEN HABE.

HEIDELBERG, DEN