

Isoforms of MUC16 activate oncogenic signaling through EGF receptors to enhance the progression of pancreatic cancer

Divya Thomas,^{1,11} Satish Sagar,^{1,11} Xiang Liu,^{1,11} Hye-Rim Lee,¹ James A. Grunkemeyer,¹ Paul M. Grandgenett,¹ Thomas Caffrey,¹ Kelly A. O'Connell,¹ Benjamin Swanson,² Lara Marcos-Silva,^{3,4} Catharina Steentoft,⁵ Hans H. Wandall,⁵ Hans Carlo Maurer,⁶ Xianlu Laura Peng,⁷ Jen Jen Yeh,⁸ Fang Qiu,⁹ Fang Yu,⁹ Ragupathy Madiyalakan,¹⁰ Kenneth P. Olive,⁶ Ulla Mandel,⁵ Henrik Clausen,⁵ Michael A. Hollingsworth,¹ and Prakash Radhakrishnan¹

¹Eppley Institute for Research in Cancer and Allied Diseases, Fred & Pamela Buffett Cancer Center, University of Nebraska Medical Center, Omaha, NE 68198-6805, USA; ²Department of Pathology and Microbiology, University of Nebraska Medical Center, Omaha, NE, USA; ³Instituto de Tecnologia Química e Biológica António Xavier, Universidade Nova de Lisboa, Av. da República, 2780-157 Oeiras, Portugal; ⁴iBET, Instituto de Biologia Experimental e Tecnológica, Apartado 12, 2780-901 Oeiras, Portugal; ⁵Copenhagen Center for Glycomics, Department of Cellular and Molecular Medicine, University of Copenhagen, Blegdamsvej 3, 2200 Copenhagen N, Denmark; ⁶Departments of Medicine and Pathology & Cell Biology, Herbert Irving Comprehensive Cancer Center, Columbia University Medical Center, New York, NY, 10032; ⁷Lineberger Comprehensive Cancer Center, University of North Carolina at Chapel Hill, Chapel Hill, NC, USA; ⁸Departments of Surgery and Pharmacology, Lineberger Comprehensive Cancer Center, University of North Carolina at Chapel Hill, Chapel Hill, NC 27514, USA; ⁹College of Public Health, Biostatistics, University of Nebraska Medical Center, Omaha, NE, USA; ¹⁰Quest PharmaTech Inc., Edmonton, Alberta T6E 6S4, Canada

Aberrant expression of CA125/MUC16 is associated with pancreatic ductal adenocarcinoma (PDAC) progression and metastasis. However, knowledge of the contribution of MUC16 to pancreatic tumorigenesis is limited. Here, we show that MUC16 expression is associated with disease progression, basal-like and squamous tumor subtypes, increased tumor metastasis, and short-term survival of PDAC patients. MUC16 enhanced tumor malignancy through the activation of AKT and GSK3 β oncogenic signaling pathways. Activation of these oncogenic signaling pathways resulted in part from increased interactions between MUC16 and epidermal growth factor (EGF)-type receptors, which were enhanced for aberrant glycoforms of MUC16. Treatment of PDAC cells with monoclonal antibody (mAb) AR9.6 significantly reduced MUC16-induced oncogenic signaling. mAb AR9.6 binds to a unique conformational epitope on MUC16, which is influenced by O-glycosylation. Additionally, treatment of PDAC tumor-bearing mice with either mAb AR9.6 alone or in combination with gemcitabine significantly reduced tumor growth and metastasis. We conclude that the aberrant expression of MUC16 enhances PDAC progression to an aggressive phenotype by modulating oncogenic signaling through ErbB receptors. Anti-MUC16 mAb AR9.6 blocks oncogenic activities and tumor growth and could be a novel immunotherapeutic agent against MUC16-mediated PDAC tumor malignancy.

INTRODUCTION

The biomarker CA125 comprises epitopes expressed on MUC16, a massive (>22,000 amino acid) membrane-bound, heavily N- and

O-glycosylated cell-surface glycoprotein.¹ Approximately 50%–70% of pancreatic ductal adenocarcinoma (PDAC) patients express CA125 and is associated with disease progression and metastasis.² In the clinical setting, high and/or increasing levels of CA125/MUC16 in circulation are widely associated with poor prognosis, whereas declining values are often associated with disease stabilization or regression.³ Antibody assays that detect CA125 also use the same capture and detection antibody, which derives from the fact that such assays detect multivalent structures that contain multiple copies of the same epitope. Unique physicochemical features of mucins include tandem repeat domains that are heavily O-glycosylated with extended structures,⁴ which explains the fact that these are the principal multivalent components that are detected by single antibody biomarker assays. Despite the prognostic importance of CA125, there is a paucity of studies into possible biological roles of MUC16 in tumor progression.

During cancer progression, some branched O-glycan structures are truncated to produce tumor-specific, short mucin-type O-glycans Tn (GalNAc alpha1-O-Ser/Thr) and sialyl Tn (Sialic acid alpha 2-6 GalNAc alpha1-O-Ser/Thr).^{5,6} Overexpression of STn antigen occurs at the highest frequency in pancreatic cancer, among other solid

Received 31 August 2020; accepted 18 December 2020;
<https://doi.org/10.1016/j.ymthe.2020.12.029>.

¹¹These authors contributed equally

Correspondence: Prakash Radhakrishnan, Eppley Institute for Research in Cancer and Allied Diseases, Fred & Pamela Buffett Cancer Center, University of Nebraska Medical Center, Omaha, NE 68198-6805, USA.

E-mail: pradhakr@unmc.edu



tumors.⁷ Also, STn expression is observed in pancreatic intraepithelial neoplasia stage III (PanIN-3), a premalignant lesion thought to precede the development of PDAC.⁸ Recent studies from our laboratory demonstrated that enforced constitutive expression of truncated O-glycans (Tn and STn antigens) on mucins and other glycoproteins through knockout (KO) of COSMC enhance the malignant potential of PDAC cells,⁹ in part through altering the steady-state signaling pathways in cells to an oncogenic phenotype. Also, we recently demonstrated that MUC16 is one of the aberrantly O-glycosylated glycoproteins detected in tumors that are expressing truncated O-glycans.¹⁰ However, the molecular and biological mechanisms by which aberrant glycoforms of MUC16 influence the pancreatic tumorigenesis are not well understood.

Here we show that MUC16 expression is clinically associated with PDAC progression, metastasis, aggressive subtypes (squamous and basal-like), and reduced patients' survival. We further investigated the biological functions of MUC16 in PDAC tumor growth by using CRISPR-Cas9 to genetically eliminate the expression of MUC16 in isogenic PDAC cells. The results show that MUC16 activated AKT and GSK3 β oncogenic signaling pathways in tumor cells through interactions with ErbB receptors. Furthermore, antibodies specific to tandem repeats of MUC16 inhibited the activation of these pathways and malignant properties of tumor cells *in vitro*, and anti-MUC16 antibody alone or in combination with gemcitabine significantly inhibited tumor growth of PDAC xenografts. The results of this study show that in addition to serving as a biomarker, MUC16 has biological activities that support the progression of pancreatic cancer.

RESULTS

MUC16 aggravates pancreatic cancer progression

Previous studies have suggested a close association between increased expression of MUC16 and PDAC progression.^{2,11} A Kaplan-Meier plot evaluated the survival probability of patients expressing low MUC16 (n = 54) and high MUC16 (n = 122) from the Human Protein Atlas revealed a significantly decreased survival probability (p = 0.0041, log-rank) in patients with higher MUC16 expression (Figure S1A). Analysis of MUC16 mRNA expression from a cohort of patients at Columbia University revealed that MUC16 was significantly elevated in PDAC epithelium and PDAC stromal cells as compared to precursor lesions PanIN-1 (FDR 1.30E-02) and IPMN (FDR 2.89E-02) epithelia. Moreover, epithelial PDAC tumors show enhanced MUC16 expression compared to stromal PDAC tumors (FDR 2.24E-47; Figure 1A; Table 1).¹² Immunohistochemical staining of a commercial pancreatic tissue microarray (BIC14011a; BIOMAX.US) containing tissues of normal pancreas and different stages of pancreatic tumors with anti-MUC16 monoclonal antibody (mAb) AR9.6 revealed significantly higher expression of MUC16 protein in ductal adenocarcinoma (DAC; p = 0.0402). Also, higher expression of MUC16 was found in adenosquamous carcinoma (ASC), and islet cell tumors compared to early PanINs (Figure 1B, representative immunohistochemistry [IHC] images; Figure 1C histoscores). Evaluation of matched sets of rapid autopsies (RAP) PDAC samples (Table

S1) by using four independent anti-MUC16 monoclonal antibodies (OC125, AR9.6, 5B9, and 5E11)^{1,13} showed high levels of MUC16 protein in primary tumors as compared to normal pancreas tissues (samples 1–7; Figure 1D, heatmap of intensity score; Figure 1E, representative IHC images). A comparison of the intensity score of MUC16 epitope detection by various MUC16 antibodies against OC125 is depicted in Figure 1F. mAb AR9.6 showed significantly high epitope detection as compared to OC125 (p < 0.0001; Table S2). About 50%–70% of PDAC patients showed MUC16 expression and detected by using different antibodies (Figure 1G). Additionally, MUC16 epitopes detected by mAb AR9.6 was increased in liver metastasis; interestingly, 5E11 was higher in primary tumors as compared to liver mets, suggesting that MUC16 expressed by primary tumors shows differential post-translational processing as compared to metastases (Figures 2A and 2B, heatmap of intensity score; Figure 2C, representative images). We also found increased expression of MUC16 in metastatic sites in addition to the liver, especially in the omentum, aorta, and appendix lesions, etc. compared to primary tumors (Figure S1B, intensity score; Figure S1C, representative IHC images). Higher expression of MUC16 is associated with primary and liver mets (Figure 2D). The percent survival of RAP patients indicated that patients with low MUC16 expression (n = 31) had survived longer than patients with high MUC16 expression (n = 30; p = 0.0257; Figure 2E). Notably, protein expression of MUC16 was significantly high (p = 0.0089) in short-term survivors (n = 34) as compared to long-term survivors (n = 32) with a median survival of 274 days (Figure 2F). Analysis of the expression of MUC16 in PDAC tumor subtype classifications from TCGA PAAD dataset¹⁴ revealed significantly increased expression of MUC16 in squamous subtype (Bailey's classifications), basal-like subtype (Moffitt classification), and quasimesenchymal subtype (Collisson classification) as compared to other subtypes (Figure 2G; Table S3). Among these, expression of MUC16 was profound in squamous subtype (n = 31) as compared to immunogenic subtype (n = 28, p < 0.001), progenitor subtype (n = 53, p < 0.001) and ADEX subtype (n = 38, p < 0.001), and compared to all the three subtypes (p = 0.0001). Also, MUC16 expression was more in the Basal-like subtype (n = 65) as compared to the classical subtype (n = 85, p < 0.001; Figure 2G).

KO of MUC16 decreases tumorigenic features of PDAC

Our previous report showed that MUC16 has an increased number of altered O-glycans in PDAC cells.¹⁰ Hence, we eliminated MUC16 in normal branched O-glycan (wild-type [WT]) and altered truncated O-glycan (SimpleCells, SC) expressing isogenic PDAC cells by using CRISPR-Cas9 deletion constructs (Figure 3A). The loss of MUC16 expression in isogenic PDAC cells (T3M4 WT-MUC16^{KO}, T3M4 SC-MUC16^{KO}, Capan-2 WT-MUC16^{KO}, and Capan-2 SC-MUC16^{KO}) was confirmed by PCR (Figures S1D–S1F), western blotting (Figures 3B and 3C), and immunofluorescence (Figure 3D). Genetic deletion of MUC16 in isogenic PDAC cells (T3M4 WT-MUC16^{KO} [2E4], T3M4 SC-MUC16^{KO} [1E10], Capan-2 WT-MUC16^{KO} [1C10], and Capan-2 SC-MUC16^{KO} [2F9]) significantly decreased cell proliferation as compared to their isogenic parental (T3M4 and Capan-2, WT and SC) cells at each

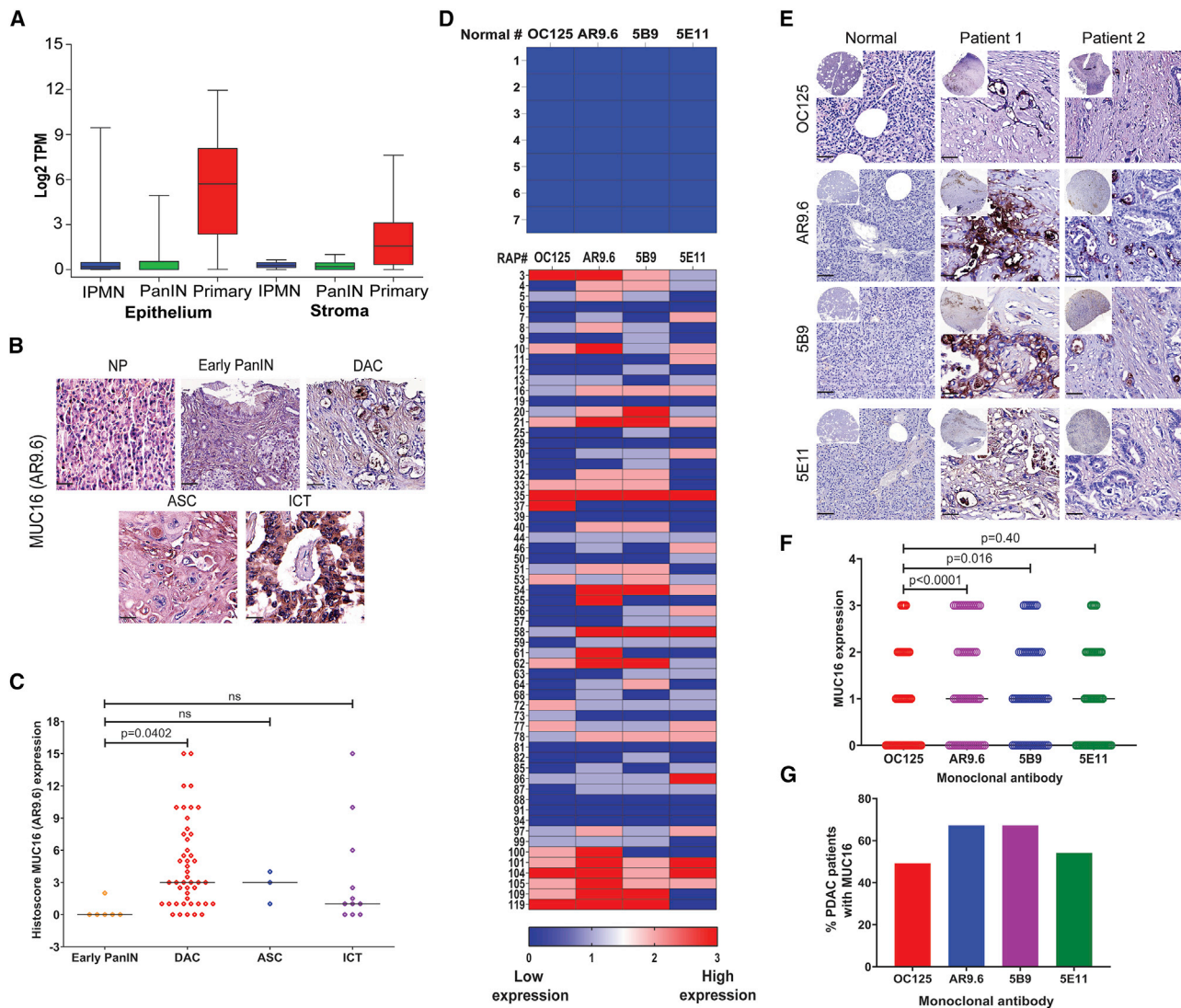


Figure 1. MUC16 in pancreatic cancer progression

(A) LCM-RNA-seq analysis of MUC16 in frozen human pancreas tissue sections (PDAC epithelium with IPMN [n = 19], PanIN [n = 26], and primary [n = 197]; and stroma with IPMN [n = 12], PanIN [n = 23], and primary [n = 124]). MUC16 mRNA expression for different samples were quantified by Log2 TPM (transcripts per million). (B) Immunohistochemical analysis of MUC16 in pancreatic tissue microarrays containing normal pancreas (NP, n = 10), early PanIN1 (n = 6), ductal adenocarcinoma (DAC, n = 46), adenosquamous carcinoma (ASC, n = 3), and islet cell tumors (n = 11) using anti-MUC16 antibody (AR9.6). Scale bar, 40 μ m. (C) Histoscore analysis of MUC16 expression by IHC. Expression of MUC16 was compared between the early PanIN and other diseased conditions. Data were presented as the median (Dunnett's multiple comparisons test). (D) Heatmap of IHC analysis of MUC16 expression (OC125, AR9.6, 5B9, and 5E11) in normal pancreatic tissues (n = 7) and RAP primary tumors (n = 61). (E) Representative IHC images of different anti-MUC16 antibodies (OC125, AR9.6, 5B9, and 5E11) stained normal pancreatic tissues and RAP primary tumors. Scale bar, 40 μ m. (F) Graphical representation of overall expression of MUC16 (OC125, AR9.6, 5B9, and 5E11) in RAP primary tumors. Data were presented as the median (n = 61) (Dunnett's multiple comparisons test). (G) Percent expression of MUC16 using different anti-MUC16 antibodies (OC125, AR9.6, 5B9, and 5E11) in RAP tumor samples.

time points ($p < 0.0001$, two-way ANOVA with Tukey's multiple comparisons; Figures 3E–3H). Cell-cycle analysis showed the accumulation of cells in the G0/G1 phase, as evidenced by reduced expression of G0/G1 phase checkpoint mediators, cyclin D1 (T3M4 and Capan-2), and cyclin E1 (T3M4) in WT- and SC-MUC16^{KO} cells (Figures 3I and 3J).

Further, we analyzed the *in vitro* and *in vivo* tumorigenic potential of isogenic PDAC cells with or without MUC16 (Figure 4A). MUC16^{KO} cells showed significantly reduced migration (T3M4 WT-MUC16^{KO}, $p = 0.0104$; T3M4 SC-MUC16^{KO}, $p < 0.0001$; Figure 4B) and invasion (T3M4 WT-MUC16^{KO}, $p = 0.0071$; T3M4 SC-MUC16^{KO}, $p < 0.0001$; Figure 4C) as compared to their parental T3M4 (WT and SC) cells.

Table 1. Differential gene-expression analysis

Compare	Where	logFC	MeanA	MeanB	t	FDR
PanIN versus PDA	Epithelium	8.36	0.30	8.65	2.97	1.30e-02
IPMN versus PDA	Epithelium	7.55	1.10	8.65	2.55	2.89e-02
PanIN versus PDA	Stroma	3.72	0.37	4.08	3.17	1.10e-02
IPMN versus PDA	Stroma	2.96	1.12	4.08	1.98	1.17e-01
Epi versus stroma	PanIN	-0.89	0.58	-0.31	-1.52	2.04e-01
Epi versus stroma	IPMN	-5.08	3.15	-1.93	-3.02	2.52e-02
Epi versus stroma	PDA	-4.45	8.80	4.35	-23.90	2.24e-47
Epi versus stroma	All	-4.41	7.73	3.32	-21.46	2.04e-49

Statistical comparison of the expression of MUC16 between different groups of samples. The statistical approach used the Voom method, which is based on linear modeling. The output includes mean TPM values for each group, the log₂ fold change (logFC) between the groups, t-statistic comparing the groups, and the False Discovery Rate (FDR) that corrects for multiple hypothesis testing.¹² Positive log-fold changes indicate higher levels in the second group.

We also detected a significantly reduced cell migration in T3M4 SC-MUC16^{KO} cells compared to T3M4 WT-MUC16^{KO} cells ($p = 0.0018$, Figure 4B). Additionally, we found a significantly reduced expression of matrix metalloproteinase 9 (MMP9) in both T3M4 WT-MUC16^{KO} and SC-MUC16^{KO} cells as compared to parental cells (Figure S1G), whereas no changes in MMP2 expression (data not shown), suggesting that MUC16 regulation of MMP9 influences migration and invasion. Consistent with previous findings,⁹ orthotopic implantation of T3M4 SC cells resulted in significantly enhanced tumor growth ($n = 14$) by weight ($p = 0.0019$, Figure 4D) and volume ($p < 0.0001$, Figure 4E) and incidence of metastases to the diaphragm (50%) and lymph node (57%), as compared to T3M4 WT (Figure 4F; Tables S4). In contrast, T3M4 SC-MUC16^{KO} cells showed significantly reduced tumor weight ($p < 0.0001$), volume ($p < 0.0001$), and metastases to peritoneum (14%, $p = 0.02$), diaphragm (7%, $p = 0.099$), lymph nodes (14%, $p = 0.05$), spleen (28%, $p = 0.0007$), and no metastasis to liver and lung as compared to T3M4 SC cells implanted tumors, (Figures 4D–4F). However, T3M4 WT-MUC16^{KO} cells did not show any significant changes as compared to T3M4 WT cells.

Aberrant glycosylation enhances the interactions between MUC16 and epidermal growth factor receptors

Ingenuity pathway analysis of truncated O-glycans expressing PDAC cells phosphoproteome dataset from our previous studies⁹ revealed that epidermal growth factor receptor family members have been activated in truncated O-glycan expressing PDAC cells (Table S5). Hence, we investigated the phosphorylation status of epidermal growth factor receptors in T3M4 WT and SC cells and found phosphorylation of ErbB1 (Y1173), ErbB3 (Y1289), and its downstream targets AKT (S473) and GSK3 β (S9) were enhanced in SC cells as compared to WT cells (Figures S2A and S2B). Evaluation of protein-protein interactions between EGF receptors (ErbB1, ErbB2, ErbB3, and ErbB4) and MUC16 using proximity ligation assay (PLA) on T3M4 WT and SC cells revealed significant interactions between MUC16 and ErbB2 (counts/cell) in WT cells that were enhanced in SC cells ($p = 0.0017$; Figure 5A). Together these data support the hypothesis that interactions between MUC16 and EGF re-

ceptors activate oncogenic signaling cascades that are exacerbated by aberrant glycosylation. Therefore, we investigated the activation of EGF type receptors and consequent downstream targets in MUC16^{KO} WT and SC PDAC cells (T3M4 and Capan-2). Consistently, there was constitutive phosphorylation of ErbB1 (Y1173), ErbB2 (Y1248), and ErbB3 (Y1289) in T3M4 and Capan-2 WT cells that was enhanced in the SC derivatives but significantly decreased or eliminated in MUC16^{KO} WT and SC clones (T3M4 and Capan-2; Figures 5B and 5C). Similarly, phosphorylation of AKT (S473) and GSK3 β (S9) was high in T3M4 and Capan-2 SC cells as compared to parental WT cells; however, there was complete inhibition of steady-state phosphorylation of AKT and GSK3 β in T3M4 SC-MUC16^{KO} cells as compared to T3M4 WT-MUC16^{KO} cells (Figure 5D). A similar trend was observed in MUC16^{KO} Capan-2 WT and SC cells (Figure 5E). These results were further confirmed in orthotopic tumors of T3M4 WT, T3M4 WT-MUC16^{KO} (2E4), T3M4 SC, and T3M4 SC-MUC16^{KO} (1E10), which showed significantly reduced expression of p-ErbB2 ($p < 0.0001$ and $p < 0.0001$; Figures 5F and 5H) and p-AKT ($p = 0.0092$ and $p = 0.0045$; Figures 5G and 5I) in T3M4 WT-MUC16^{KO} and T3M4 SC-MUC16^{KO} tumors as compared to control tumors, respectively. Interestingly, expression of p-ErbB2 was significantly less in T3M4 SC-MUC16^{KO} tumors ($p = 0.0068$) compared to T3M4 WT-MUC16^{KO} tumors. To confirm the clinical relevance and the posited regulatory role of MUC16 on EGFR signaling, we purified MUC16/CA125 from PDAC patients ($n = 5$) and applied this to T3M4 WT-MUC16^{KO} cells (5 μ g/mL for 24 h). We observed increased phosphorylation of ErbB3, ErbB2, AKT, and GSK3 β in clinically isolated MUC16 treated MUC16^{KO} cells compared to untreated cells (Figure S2C).

Anti-MUC16 antibodies treatment inhibits EGF receptors and their downstream signaling

We further evaluated the contribution of MUC16 to the constitutive activation of EGF receptors and downstream oncogenic signaling by using different anti-MUC16 mAbs to block MUC16 binding to receptors on tumor cells. We have previously reported the binding specificity of mAbs 5E11¹³ and B43.13 to MUC16.¹⁵ Treatment of T3M4 WT cells with anti-MUC16 antibodies (5E11, B43.13, and AR9.6;

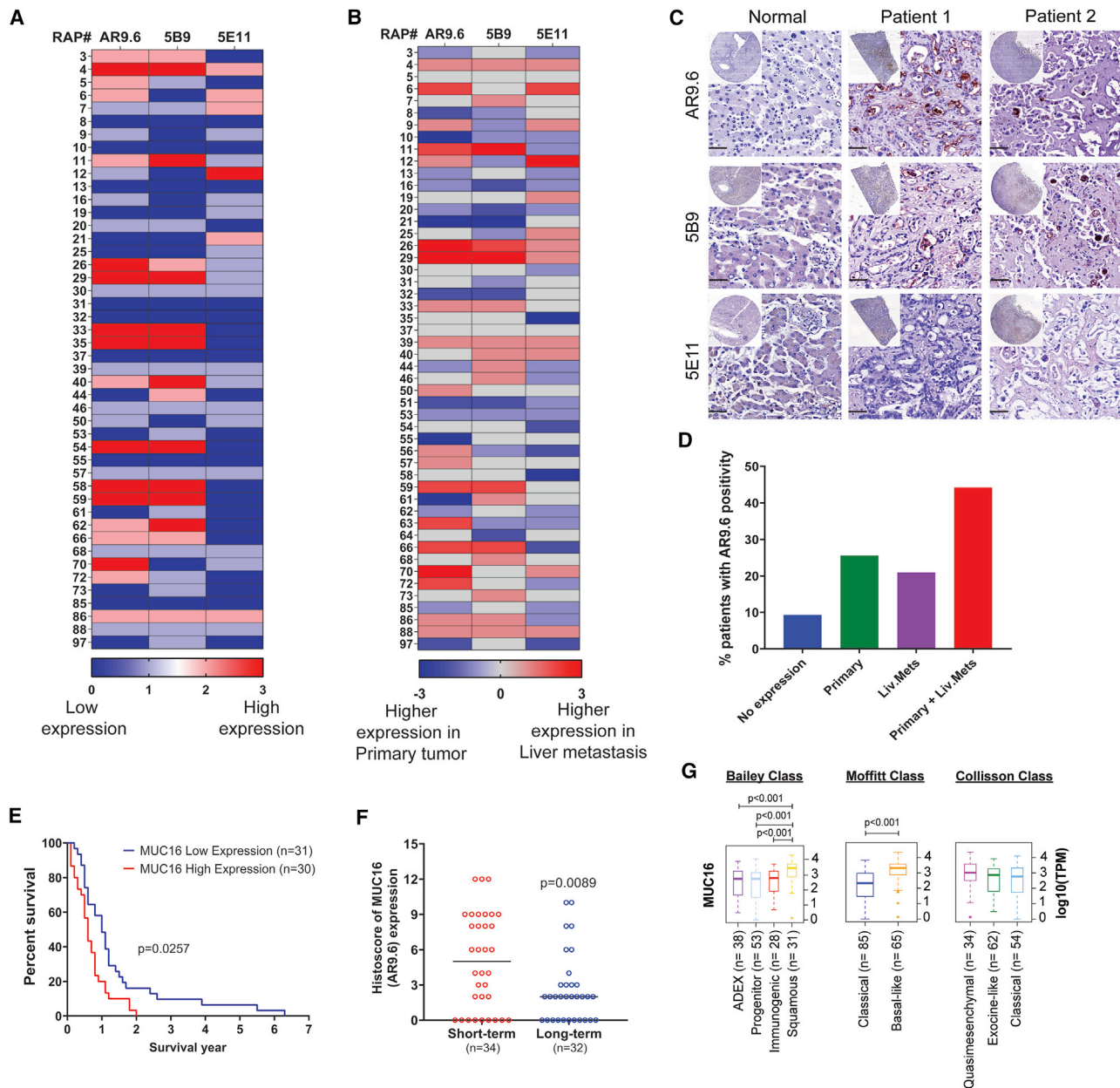


Figure 2. MUC16 in pancreatic tumor metastasis

(A) Heatmap of IHC analysis of MUC16 expression (AR9.6, 5B9, and 5E11) in liver metastasis (n = 46). (B) Heatmap of comparison of MUC16 expression in matched sets of RAP primary tumors and liver mets. Higher intensity of color corresponds to high expression of MUC16 based on immunohistochemical score. (C) Representative images of IHC analysis of MUC16 expression in normal liver tissue and RAP liver mets using MUC16-specific antibodies AR 9.6, 5B9, and 5E11. Scale bar, 40 μ m. (D) Percent expression of MUC16 (AR9.6) in RAP tumor tissues of primary tumor alone, liver mets alone, and primary tumor and liver mets together. (E) Survival of RAP cohort patients with high and low MUC16 (AR9.6) expression (Mantel-Cox test). (F) Histoscores of MUC16 (AR9.6) compared between short-term (n = 34) and long-term (n = 32) PDAC survivors. Data were presented as median (unpaired t test). (G) Boxplot of MUC16 expression level stratified by RNA expression dataset from Bailey class, Moffitt class, and Collisson class of PDAC. Boxplots were generated by comparing the expression of MUC16 gene among the subtypes using Mann-Whitney rank-sum test. A p value of less than 0.01 was considered statistically significant.

2.5 and 5 μ g/mL) for 24 h inhibited the basal level phosphorylation of AKT in a concentration-dependent manner (Figure 6A). The greatest inhibition of AKT phosphorylation was observed in cells treated with

mAb AR9.6 (2.5 and 5 μ g/mL). Among these antibodies, only mAb AR9.6 decreased constitutively hyperphosphorylated AKT in T3M4 SC cells compared to untreated and immunoglobulin G (IgG) control

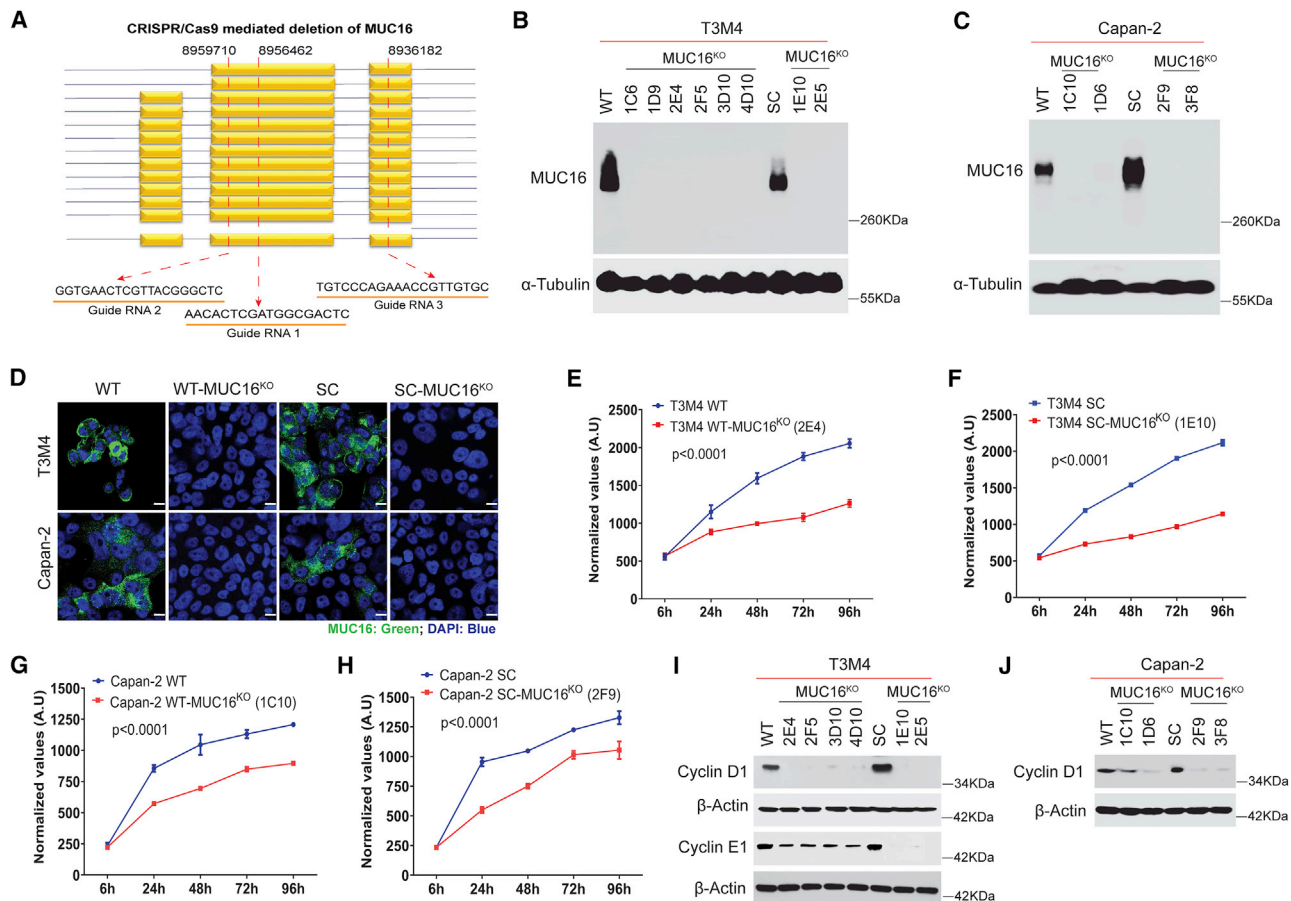


Figure 3. Genetic deletion of MUC16 in PDAC cells

(A) Schematic representation of the targeted deletion of MUC16 via CRISPR-Cas9 constructs in T3M4 and Capan-2 (WT and SC) cells with three gRNA target loci are shown as red dotted lines. (B and C) Western blotting of MUC16 in T3M4 and Capan-2 (WT and SC) parental cells and MUC16^{KO} clones respectively. Detection of α -tubulin served as a loading control. (D) Immunofluorescence analysis of MUC16 in T3M4 WT, WT-MUC16^{KO} (2E4), T3M4 SC, SC-MUC16^{KO} (1E10), Capan-2 WT, WT-MUC16^{KO} (1C10), Capan-2 SC, and SC-MUC16^{KO} (2F9) cells. Scale bar, 10 μ m. (E–H) Cell proliferation assays in T3M4 WT, WT-MUC16^{KO} (2E4) (E), T3M4 SC, SC-MUC16^{KO} (1E10) (F), Capan-2 WT, WT-MUC16^{KO} (1C10) (G), and Capan-2 SC, SC-MUC16^{KO} (2F9) (H) cells. Data were presented as mean \pm SD (n = 6; Tukey's multiple comparisons test). (I) Western blotting of Cyclin D1 and Cyclin E1 in T3M4 WT, WT-MUC16^{KO} clones, SC, and SC-MUC16^{KO} clones. (J) Western blotting of Cyclin D1 in Capan-2 WT, WT-MUC16^{KO} clones, SC, and SC-MUC16^{KO} clones. β -actin was used as a loading control.

(Figure 6B). Next, we analyzed the phosphorylation of AKT upstream activator EGF receptors in mAb AR9.6-treated PDAC cells and found reduced phosphorylation of ErbB1 and ErbB2 (Figure S3A). These results prompted us to characterize the MUC16 epitope recognized by mAb AR9.6. An established epitope-mapping strategy was used that utilizes nine MUC16 constructs with overlapping sequences within the tandem repeat (TR) region of MUC16 (TR1.7), includes part of TR4, the entire Sea urchin, enterokinase and agrin (SEA), and linker domain of TR5 along with half of the SEA domain of TR6. The TR1.2, 1/4, 5/2, 5/7, 5/8, 5/9, K292, and F/R constructs represent truncated forms of this construct (Figure 6C).¹ ELISA assay of MUC16 constructs that were expressed in *E. coli* and purified by nickel chromatography with mAb AR9.6 showed strong reactivity of mAb AR9.6 with tandem repeats TR1.7 and TR1.2. In contrast, tandem repeats 5/2, 1/4, 5/7, 5/8, 5/9, K292, F/R were unreactive (Figure 6D). This

distinguishes mAb AR9.6 from the classical OC125 and M11 epitopes that were strongly reactive with the 5/2 construct. Also, incubation of proteins with anti-MUC16 mAb 5E11 served as a positive control for these reactions, which showed reactivity with proteins encoded by the TR1.7, TR1.2, 5/2, 1/4, and 5/7 constructs (Figure 6E). The incubation of proteins with a negative control showed no reactivity toward the above-mentioned TR constructs (data not shown). We found that mAb AR9.6 binds to multiple isoforms of MUC16 that are distinct from those bound by mAbs M11 and OC125. Additionally, we found that mAb 5E11 binds to a distinct isoform of MUC16 as compared to other anti-MUC16 mAbs (Figure S3B).

Next, we investigated the possible biological activity of the AR9.6 binding region on MUC16 (TR1.2) by expressing and purifying MUC16 TR1.2 from CHO cells (Figure S3C) and treating T3M4-

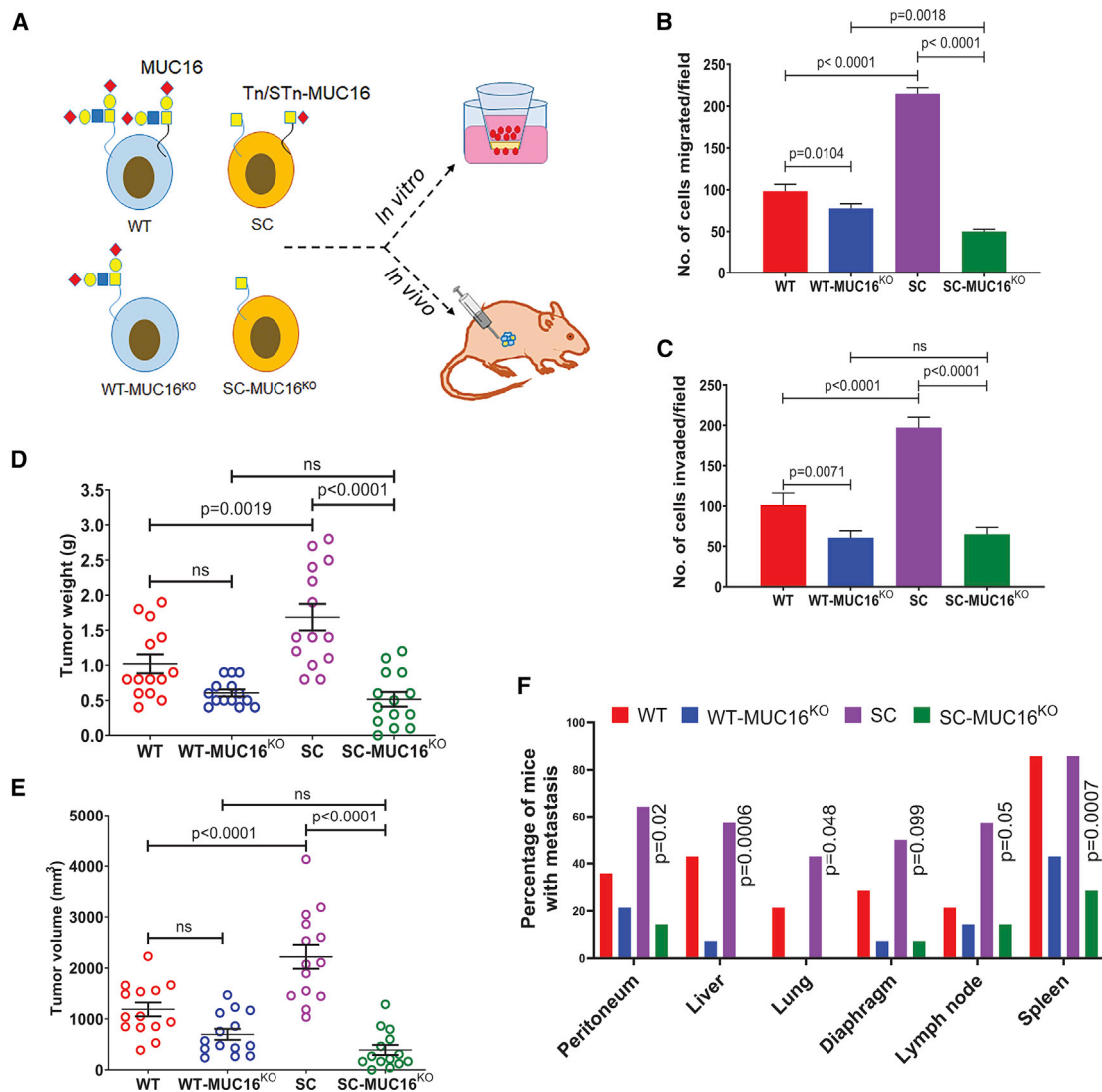


Figure 4. Genetic deletion of MUC16 reduces PDAC tumorigenicity

(A) Schema of *in vitro* and *in vivo* experiments using WT, WT-MUC16^{KO}, SC, and SC-MUC16^{KO} cells. (B) Cell migration assay in T3M4 WT, WT-MUC16^{KO} (2E4), T3M4 SC, and SC-MUC16^{KO} (1E10) cells. Data were presented as mean ± SD (n = 3; Dunnett's multiple comparisons test). (C) Matrigel invasion assay in T3M4 WT, WT-MUC16^{KO} (2E4), T3M4 SC, and SC-MUC16^{KO} (1E10) cells. Data were presented as mean ± SD (n = 3; Dunnett's multiple comparisons test). (D and E) Tumor weight (D) and tumor volume (E) of T3M4 WT, WT-MUC16^{KO} (2E4), T3M4 SC, and SC-MUC16^{KO} (1E10) cells implanted orthotopic tumors. Data were presented as mean ± SEM (n = 14; Dunnett's multiple comparisons test). (F) Analysis of tumor metastasis in T3M4 WT, WT-MUC16^{KO} (2E4), T3M4 SC, and SC-MUC16^{KO} (1E10) cells implanted tumor-bearing animals (Fisher's exact test).

MUC16^{KO} cells. There were dose-dependent increases in the phosphorylation of ErbB1, ErbB3, and GSK3β upon the addition of TR1.2 to MUC16 deficient PDAC cells (Figure 6F). Pre-incubation of MUC16^{KO} PDAC cells with mAb AR9.6 significantly inhibited TR1.2 induced phosphorylation of p-ErbB3, p-ErbB2, p-AKT, and p-GSK3β (Figure 6G). We investigated the role of glycosylation in TR1.2 binding affinity with mAb AR9.6. Treatment of TR1.2 with N-glycanase resulted in reduced mobility and binding to mAb AR9.6; however, treatment of TR1.2 with either Sialidase or O-glycanase or both significantly enhanced the binding of mAb AR9.6 with

TR1.2 (Figure S3D). A similar binding trend was observed when glycosidases were applied to MUC16-positive PDAC cells (Figure S3E).

mAb AR9.6 treatment reduces *in vivo* tumor growth and metastasis

Next, we evaluated the preclinical *in vivo* therapeutic efficacy of mAb AR9.6 in an orthotopic pancreas tumor model system. mAb AR9.6-treated T3M4 WT tumor-bearing animals showed a significantly reduced tumor growth (by tumor weight, g; p = 0.01; Figure 6H), metastases to the spleen (15.34%, p = 0.04) and lymph nodes (7.69%, p =

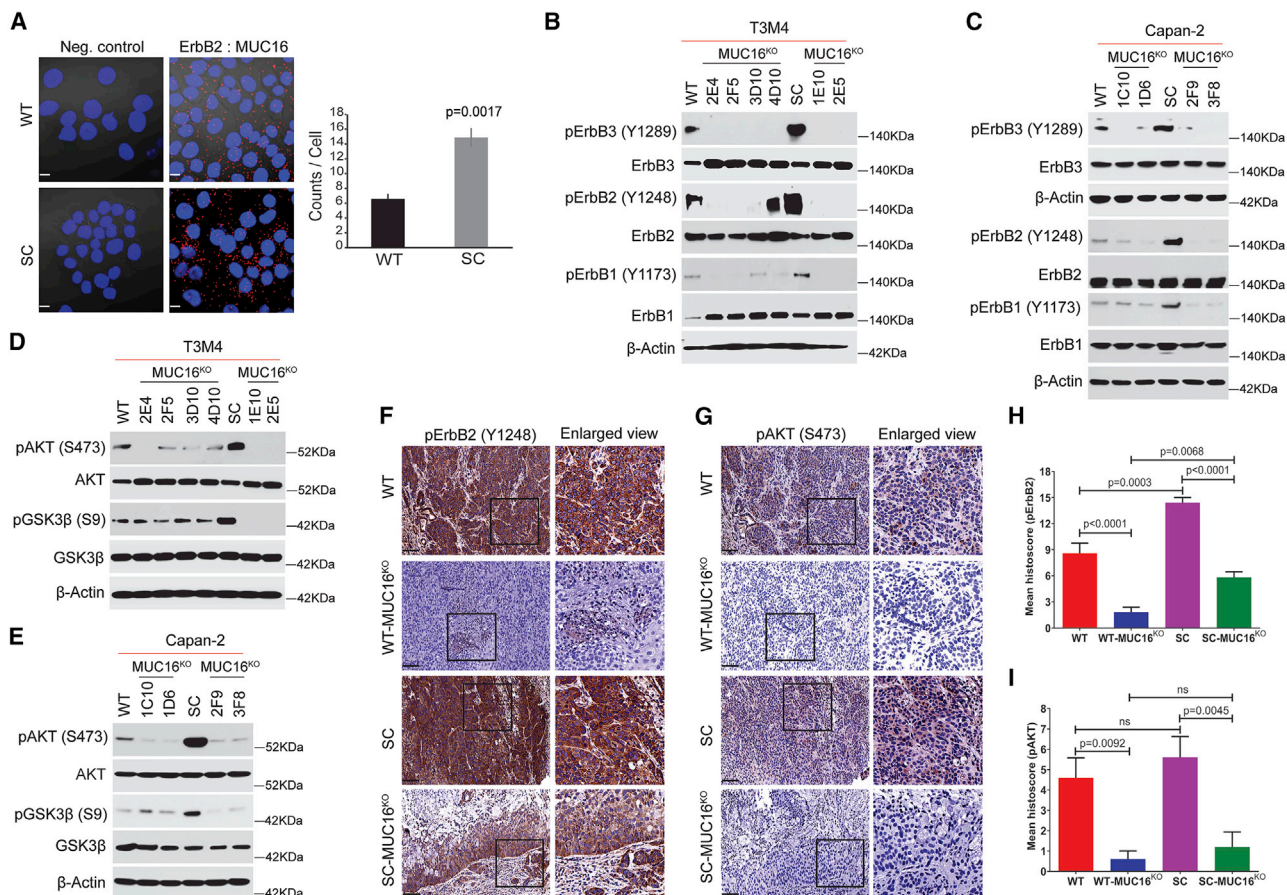


Figure 5. Interaction of MUC16 with epidermal growth factor receptors

(A) Proximity ligation assay in T3M4 WT and SC cells using MUC16 and ErbB2 specific antibodies. Scale bar, 10 μ m. Quantification of the number of interactions (dots/cell) quantified by blob finder 2 software. Cells incubated with a single antigen-specific antibody served as a negative control. Data were presented as mean \pm SD ($n = 3$; unpaired t test). (B) Western blotting of T3M4 WT, WT-MUC16^{KO}, T3M4 SC, and SC-MUC16^{KO} cell lysates with p-ErbB3 (Y1289), ErbB3, p-ErbB2 (Y1248), ErbB2, p-ErbB1 (Y1173), and ErbB1. (C) Western blotting of Capan-2 WT, WT-MUC16^{KO}, Capan-2 SC, and SC-MUC16^{KO} cell lysates with p-ErbB3 (Y1289), ErbB3, p-ErbB2 (Y1248), ErbB2, p-ErbB1 (Y1173), and ErbB1. (D) Western blotting of p-AKT (S473), AKT, p-GSK3 β (S9), and GSK3 β in T3M4 WT, WT-MUC16^{KO}, T3M4 SC, and SC-MUC16^{KO} cells. (E) Western blotting of p-AKT (S473), AKT, p-GSK3 β (S9), and GSK3 β in Capan-2 WT, WT-MUC16^{KO}, Capan-2 SC, and SC-MUC16^{KO} cells. Detection of β -actin served as a loading control. (F and G) Immunohistochemical analysis of p-ErbB2 (F) and p-AKT (G) in T3M4 WT, WT-MUC16^{KO} (2E4), T3M4 SC, and SC-MUC16^{KO} (1E10) cells implanted mouse tumors ($n = 5$). Scale bar, 40 μ m. (H and I) Mean histoscore of the IHC analysis of p-ErbB2 (H) and p-AKT (I) in T3M4 WT, WT-MUC16^{KO} (2E4), T3M4 SC, and SC-MUC16^{KO} (1E10) cells implanted mouse tumors. Data were presented as mean \pm SEM ($n = 5$; Dunnett's multiple comparisons test).

0.003), and no metastasis to lung and diaphragm as compared to vehicle control-treated T3M4 WT tumor-bearing animals (Figure 6I). We did not observe detectable changes in tumor weight (g) between the different treatment conditions on T3M4 SC tumor-bearing animals (vehicle control and mAb AR9.6; Figure S3F). However, a significantly reduced tumor metastasis to the peritoneum ($p < 0.0001$), no metastasis to lung, spleen, and diaphragm, and a lower percent to the liver and lymph node were found in mAb AR9.6-treated T3M4 SC tumor-bearing animals as compared to vehicle control (Figure 6J). This finding was of great interest that mAb AR9.6 significantly reduces metastasis in tumor-bearing animals. To scrutinize the mechanism behind this, we have analyzed the expression of MMP9 in AR9.6 treated T3M4 cells. We detected a reduced expression of MMP9 in mAb AR9.6-treated PDAC cells (Figure S3G).

mAb AR9.6 and gemcitabine combination therapy reduces PDAC tumorigenesis

Since we found a significant reduction in the tumor growth and metastasis upon treatment with mAb AR9.6, we sought to analyze the potential of combination therapy of AR9.6 with gemcitabine (GEM) as it is a standard chemotherapeutic agent for PDAC. Prior to the *in vivo* experiments, we assessed the potential of AR9.6 + GEM in inducing *in vitro* cell death in PDAC cells (T3M4 and Capan-2) using live/dead cytotoxicity assay (Figure 6K). The combination of AR9.6 + GEM significantly induced cell death as compared to vehicle control ($p < 0.0001$ and $p < 0.0001$), GEM alone ($p < 0.0001$ and $p < 0.0001$), and AR9.6 alone ($p < 0.0001$ and $p = 0.0001$) in both T3M4 and Capan-2 cells, respectively (Figures 6L and 6M). To further evaluate its *in vivo* therapeutic potential, we treated T3M4 tumor-

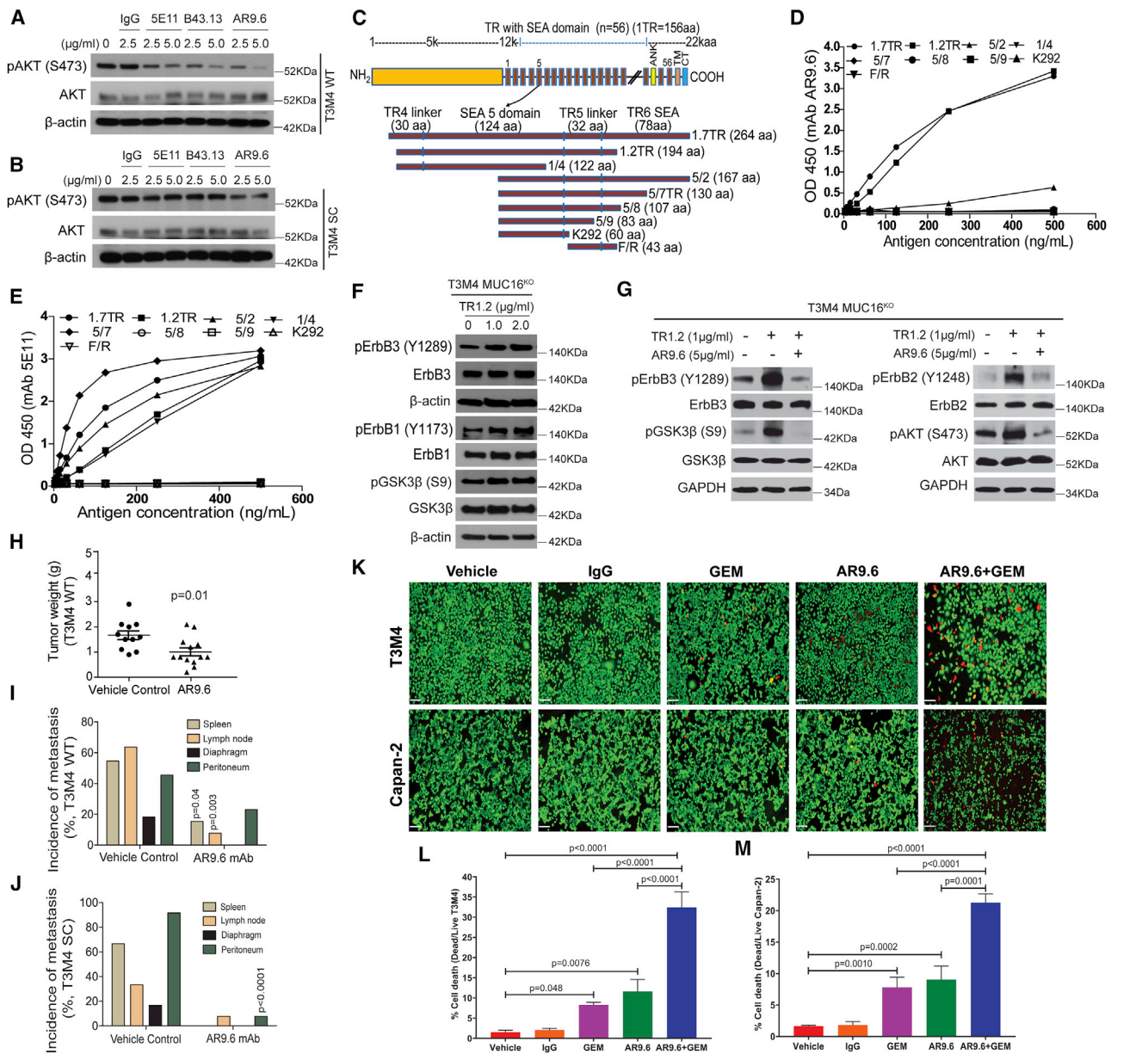


Figure 6. AR9.6 mAb inhibits MUC16 induced oncogenic signaling

(A and B) Western blotting of p-AKT and AKT in T3M4 WT (A) and T3M4 SC (B) cells treated with IgG and MUC-16 specific antibodies 5E11, B43.13, and AR9.6. Detection of β -actin served as a loading control. (C) Graphic representation of MUC16 tandem repeats constructs expressed in *E. coli*. TR1.7 includes part of TR4, the entire SEA, and the linker domain of TR5 along with half of the SEA domain of TR6 (12,660–12,923 aa). The TR1.2 (12,665–12,858 aa), 1/4 (12,665–12,785 aa), 5/2 (12,757–12,923 aa), 5/7 (12,757–12,886 aa), 5/8 (12,757–12,863 aa), 5/9 (12,757–12,839 aa), K292 (12,757–12,816 aa), and F/R (12,817–12,859 aa) constructs represent truncated forms of this construct. (D) mAb AR 9.6 epitope characterization by ELISA assay of MUC16 constructs. (E) ELISA assay of MUC16 constructs with positive control mAb 5E11. (F) Western blotting of p-ErbB3 (Y1289), ErbB3, p-ErbB1 (Y1173), ErbB1, p-GSK3 β (S9), and GSK3 β in MUC16^{KO} T3M4 cells treated with TR1.2 (1 and 2 μ g/mL, 14 h). (G) Western blotting of p-ErbB3 (Y1289), ErbB3, p-ErbB2 (Y1248), ErbB2, p-AKT (S473), AKT, p-GSK3 β (S9), and GSK3 β in MUC16^{KO} T3M4 cells treated with TR1.2 (1 μ g/mL) with or without mAb AR9.6 (5 μ g/mL). Detection of β -actin and GAPDH served as the loading control. (H) Tumor weight of vehicle control (n = 13) and mAb AR9.6 (n = 13) treated T3M4 WT orthotopic tumors. Data were reported as mean \pm SEM (unpaired t test). (I and J) Analysis of tumor metastasis in vehicle and mAb AR9.6 treated T3M4 WT (I) and SC (J) orthotopic tumor-bearing animals (Fisher's exact test). (K) Representative images of live/dead cytotoxicity assay in vehicle, IgG, AR9.6, GEM, and AR9.6 + GEM treated PDAC (T3M4 and Capan-2) cells. Scale bar, 20 μ m. (L and M) The percentage of cell death in vehicle, IgG, AR9.6, GEM, and AR9.6 + GEM-treated T3M4 (L), and Capan-2 (M) cells. Data presented as mean \pm SD (n = 3; Dunnett's multiple comparisons test).

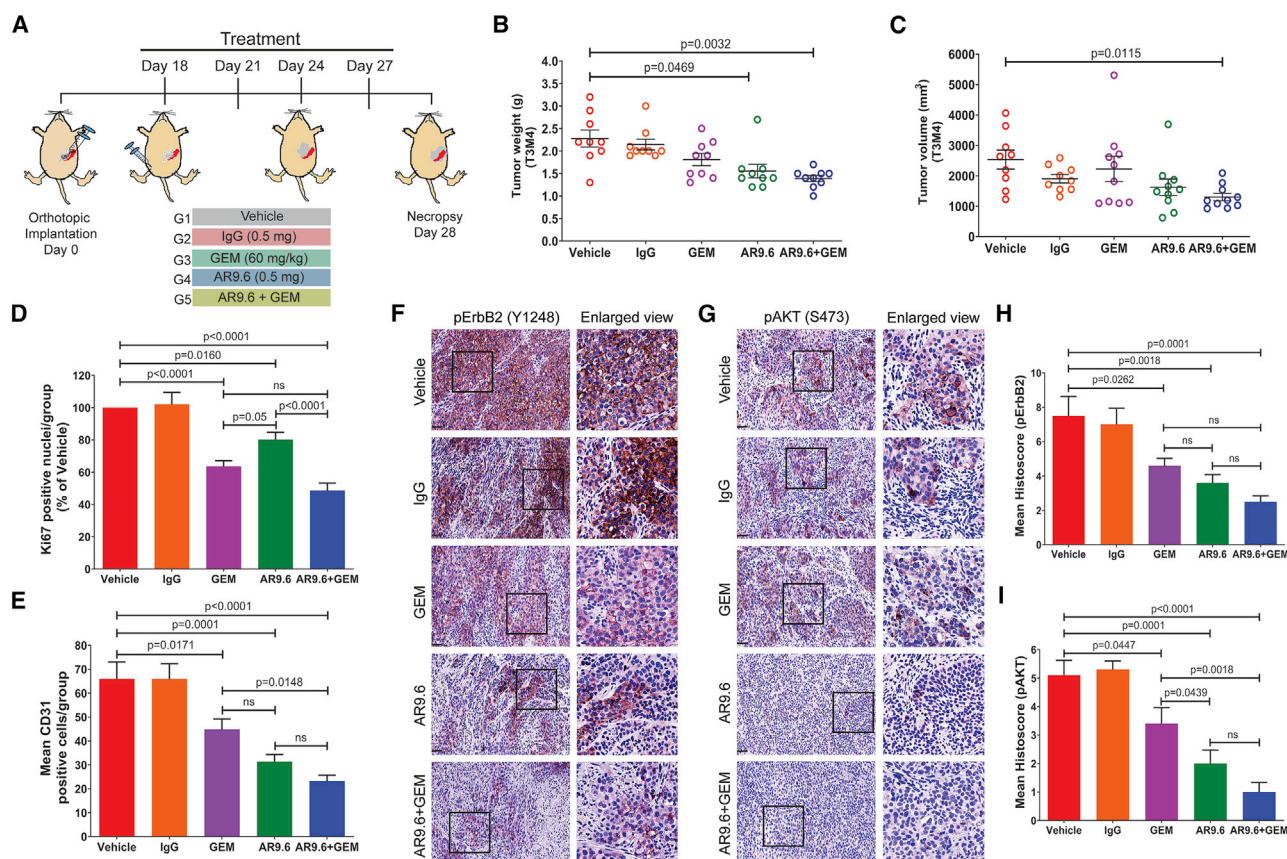


Figure 7. mAb AR9.6 plus gemcitabine treatment reduces *in vivo* tumor growth

(A) Schematic representation of orthotopic tumor implantation and treatment schedules of tumor-bearing mice ($n = 10$) with PBS, IgG, GEM, mAb AR9.6, and mAb AR9.6 plus GEM. (B and C) Tumor weight (g) (B) and tumor volume (mm^3) (C) of vehicle, IgG, GEM, mAb AR9.6, and GEM + mAb AR9.6-treated T3M4 orthotopic tumors. Data presented as mean \pm SEM ($n = 10$; Dunnett's multiple comparisons test). (D and E) IHC analysis of Ki-67 (D) and CD31 (E) in above mentioned experimental tumors. Data presented as mean \pm SEM ($n = 9$; Dunnett's multiple comparisons test). (F and G) IHC analysis of p-ErbB2 (F) and p-AKT (G) in above mentioned experimental tumors. (H and I) Mean histoscore of the IHC analysis of p-ErbB2 (H) and p-AKT (I) in above mentioned experimental tumors. All treatment groups were compared against vehicle control (PBS) treated groups. Scale bar, 40 μm . Data presented as mean \pm SEM ($n = 9$; Dunnett's multiple comparisons test).

bearing athymic nude mice with AR9.6 + GEM (Figure 7A). Since we found that mAb AR9.6 significantly reduced tumor growth in T3M4 WT tumor (Figure 6H), we sought to analyze its effect in the later stage of tumor growth. Therefore, we treated tumor-bearing mice with mAb AR9.6, GEM or AR9.6 + GEM after 18 days of PDAC cells implantation. mAb AR9.6 + GEM combination therapy produced a maximum reduction in the tumor weight ($p = 0.0032$) and volume ($p = 0.0115$) when compared to the vehicle control group (Figures 7B and 7C). Also, animals treated with mAb AR9.6 alone showed a significant reduction in tumor weight ($p = 0.0469$) when compared to vehicle control (Figure 7B) even at the late time point of treatment. While there was a reduction in tumor growth, animals treated with GEM alone did not exhibit a significant reduction in tumor weight and volume probably due to the treatment at later stage of tumor growth. Treatment of tumor-bearing animals with mAb AR9.6 and GEM significantly reduced Ki-67 ($p < 0.0001$; Figure 7D, representative positive nuclei percentage of vehicle control; Figure S4A, representative IHC images) and CD31 ($p < 0.0001$) expression (Figure 7E, positive

representative cells; Figure S4B, representative IHC images). Notably, the phosphorylation of ErbB2 and AKT was significantly inhibited in tumors from animals treated with mAb AR9.6 + GEM (p-ErbB2, $p = 0.0001$; p-AKT, $p < 0.0001$; Figures 7F and 7G representative IHC images; and Figures 7H and 7I, representative mean histoscore). Also, a significant reduction in p-ErbB2 and p-AKT expression was observed in AR9.6 alone or gemcitabine alone treated tumors as compared to vehicle control tumors (Figures 7F–7I).

DISCUSSION

MUC16 is not expressed in the normal pancreas, and aberrant expression of MUC16 is associated with pancreatic cancer progression, metastasis, and poor prognosis. Here, we show that aberrant expression of MUC16 is related to epithelial tumors, squamous, basal-like, and quasimesenchymal subtypes of PDAC, which are generally associated with an aggressive phenotype. Aberrant expression of MUC16 was also found in pancreatic adenosquamous and islet cell tumors. We found that short-term survivors of PDAC possess a higher

expression of MUC16, which is in accordance with the previous reports.¹⁶ However, the pathological mechanism of MUC16 in PDAC progression and metastasis is poorly understood. Here, we observed that interactions between MUC16 and ErbB receptors facilitate PDAC progression through the activation of AKT/GSK3 β oncogenic signaling that was further enhanced for aberrant glycoforms of MUC16. Genetic deletion of MUC16 in WT and to a greater extent in truncated O-glycan expressing PDAC cells significantly inhibited the activation of ErbB receptors and downstream oncogenic signaling pathways, further supporting the hypothesis that EGF-like domains on MUC16 bind and activate ErbB type receptors in a glycosylation modulated manner. As well, the genetic deletion of MUC16 significantly reduced tumor growth and metastasis (especially to the peritoneum and lymph node). These findings extend our previous report that MUC16 is upregulated in metastatic PDAC² to show that MUC16 has pro-oncogenic activities and supports findings from other groups showing that aberrant expression of MUC16 facilitates peritoneal metastasis.¹⁷

Activation of ErbB receptors and their downstream effectors PI3K/AKT/GSK3 β affect cellular processes, including cell proliferation, growth, motility, and cell survival.¹⁸ More interestingly, the squamous subtype of PDAC was shown to be associated with activation of EGF receptor and PI3K/AKT signaling cascades.¹⁹ More than 83% of the pancreatic tumors show constitutively activated AKT (S473), which portends poor prognosis.^{20–22} Increased expression of MUC16 in the squamous subtype of PDAC tumors along with the genetic deletion of MUC16 decreased steady-state activation of ErbB, AKT, and GSK3 β oncogenic signaling strongly support our hypothesis that MUC16 promote PDAC tumorigenicity through the interaction with ErbB receptors. However, the functional consequences of normal and altered O-glycosylation of MUC16 during PDAC progression remain elusive. Generally, glycosylation can block potential sites of other post-translational modifications of proteins,^{23,24} and glycosylated moieties may bind directly to lectin-like motifs to facilitate or block the formation of protein complexes. We found that the association between MUC16 and ErbBs in pancreatic cancer is highly influenced by glycosylation. The results of this study support the hypothesis that normal and aberrant glycosylation influences the accessibility of domains on mucins (MUC16) that bind to and/or otherwise activate cell-surface ErbB-type receptors and consequent signaling programs, which in turn affects the malignant properties of PDAC cells. It is a well-known fact that aberrant glycoforms of MUC16 is a biomarker that is elevated in the circulation in several adenocarcinomas, and its high levels are associated with poor prognosis. Treatment of MUC16 null cells with purified MUC16 from PDAC patients' ascites fluid induces an increased expression of phosphorylated EGF receptors and its downstream target signaling effectors, which supports our hypothesis that differential glycosylation of this biomarker creates active compounds that can bind different receptor tyrosine kinases and that these glycoforms could enhance disease progression by binding and activating signaling cascades in the local tumor microenvironment and at distant organ sites following circulatory transit.

The observed oncogenic activity of MUC16 in PDAC biology suggests a need for the development of therapeutic agents that target this tumor biomarker. In this context, we have characterized mAb AR9.6 that binds specifically to SEA domain 5 of MUC16. Binding of mAb AR9.6 to MUC16 inhibited phosphorylation of ErbB-type receptors and downstream AKT/GSK3 β signaling. mAb AR9.6 was found to inhibit T3M4 WT cells implanted PDAC tumor growth and metastasis in athymic nude mice alone or in combination with gemcitabine. However, mAb AR9.6 did not affect tumor growth in T3M4 SC cells implanted athymic nude. This can be explained by the abundance of MUC16 epitope (opening of hidden epitopes) on SC cells, which may need more concentration of antibody to block the epitope in order to reduce tumor growth that needs further evaluation. AR9.6 mAb staining on clinical specimens of PDAC tumors revealed that MUC16 is not only expressed in ductal adenocarcinoma but also highly expressed in adenosquamous carcinoma and islet cell tumor. Additionally, mAb AR9.6 binding epitopes were identified in a greater number of patient samples than OC125 mAb epitopes. Further investigation of AR9.6 and other MUC16 reactive antibodies as potential therapeutic agents is warranted.

MATERIALS AND METHODS

Human rapid autopsy samples

De-identified sets of samples were procured from patients who underwent a Rapid Autopsy Program at the University of Nebraska Medical Center (UNMC) with proper consent and approved by Institutional Review Board (IRB 091-01-EP). A primary pancreatic tumor (n = 72), metastatic tumors (n = 46), and normal pancreas tissues (n = 7) were used for the study (Table S2).

Cell lines and culture

Human PDAC cells T3M4 and Capan-2 (WT [do not express truncated O-glycans]; COSMC KO, SC do express truncated O-glycans)^{9,10,25} were maintained in Dulbecco's modified Eagle's medium (DMEM) and RPMI medium, respectively (Hyclone, Logan, UT, USA) supplemented with 10% fetal bovine serum (FBS; Valley Biomedical, Winchester, VA) and 100 units/mL penicillin and 100 μ g/mL streptomycin (Mediatech, Manassas, VA, USA). CHO cells were cultured in 80% EX CELL CHO Cloning Medium and 20% EX CELL CD CHO Fusion medium (Sigma, St. Louis, MO, USA) with 2% glutamine and 0.32 mg/mL G418 (Invitrogen, Carlsbad, CA, USA). All the cell lines were maintained at 37°C with 5% CO₂ in a humidified incubator.

Genetic deletion of MUC16 in PDAC cells

The N-terminal region of the MUC16 gene was genetically deleted in T3M4 and Capan-2 (WT and SC) cells by using a pre-made CRISPR-Cas9 KO plasmid kits (Santa Cruz Biotechnology, CA, USA) as per the manufacturer's instruction. Briefly, cells were transfected with a pool of three MUC16 guide-RNA-containing vectors (1, 5'-AACACACTCGATGGCGACTC-3'; 2, 5'-GGTGAACCTCGT-TACGGGCTC-3'; 3, 5'-TGTCCCAGAAACCGTTGTGC-3'), which targets MUC16 genome. MUC16 KO cells (T3M4 WT-MUC16^{KO},

T3M4 SC-MUC16^{KO}, Capan-2 WT-MUC16^{KO}, and Capan-2 SC-MUC16^{KO} were utilized for further studies.

Cell proliferation assay

Cell proliferation assay was performed as described.²⁶ Briefly, 1×10^3 PDAC cells (T3M4 and Capan-2) WT, WT-MUC16^{KO} (2E4 and 1C10), SC, and SC-MUC16^{KO} (1E10 and 2F9), respectively, were seeded into 96-well plates in quadruplicate and incubated for 24, 48, 72, and 96 h, respectively. At the end of each time, 20 μ L of cell suspension was removed from each well, and 20 μ L Alamar blue reagent (Invitrogen, Carlsbad, CA, USA) was added, such that Alamar blue made up 10% of the total volume of each well and incubated for further 2–4 h at 37°C. The fluorescence intensity was measured using Spectramax M5e (Molecular Devices, Sunnyvale, CA, USA) spectrofluorimeter at excitation 540 nm and emission 590 nm.

Cell migration assay

In vitro cancer cell migration assay was performed as described previously.²⁶ Briefly, T3M4 WT, WT-MUC16^{KO} (2E4), SC, and SC-MUC16^{KO} (1E10) cells (5×10^4 in serum-free medium) were seeded on the top of the polyethylene terephthalate membrane (PET; 24-well insert, 8 μ m; BD Biosciences, Bedford, MA, USA). After 18 h of incubation, cells that didn't migrate through the membrane were removed with cotton swabs, and cells that migrated through the membrane were fixed and stained with Diff-Quick cell staining kit (Hema3 stat pack, Thermo Fisher Scientific, Waltham, MA, USA), and counted under a light microscope in 4 different random fields at 20 \times magnification.

Cell invasion assay

Tumor cell invasion assay was performed as described previously.²⁶ In brief, T3M4 WT, WT-MUC16^{KO} (2E4), SC, and SC-MUC16^{KO} (1E10) cells (5×10^4 in serum-free medium) were seeded on Matrigel-coated membranes (BD Biosciences, Bedford, MA, USA) and incubated for 30 h at 37°C. After incubation, non-invading cells on the upper surface of the filter were removed with cotton swabs, and cells that invaded through the Matrigel were fixed and stained with Diff-Quick cell stain kit (Hema3 stat pack, Thermo Fisher Scientific, USA). The invaded cells were counted and analyzed.

Immunofluorescence analysis

Immunofluorescence staining of mucins and mucin glycans on cancer cells was carried out as described previously.⁹ Briefly, after fixing with 4% paraformaldehyde and permeabilized with 0.15% Triton X-100 in 1X PBS containing 1% BSA, PDAC cells (T3M4 and Capan-2) WT, WT-MUC16^{KO} (2E4 and 1C10), SC, and SC-MUC16^{KO} (1E10 and 2F9), respectively, were incubated with MUC16-specific mAb AR9.6, and control mouse IgG for 2 h at room temperature. Cells were washed and incubated with Alexa Fluor 488-conjugated goat anti-mouse IgG secondary antibodies (Jackson ImmunoResearch, West Grove, PA, USA). Cells were washed and mounted with Vectashield mounting medium with 4',6-diamidino-2-phenylindole (DAPI) (Vector Laboratories, Burlingame, CA, USA). Immunofluo-

rescence images were captured by using the Zeiss LSM 710 confocal laser scanning microscope (Carl Zeiss, Thornwood, NY, USA; Confocal Laser Scanning Fluorescence Microscopy Core Facility, UNMC).

Orthotopic pancreas tumor model

All the animals were housed under standard housing conditions at the University of Nebraska Medical Center animal core facilities, and all animal procedures were reviewed and approved by the UNMC institutional animal care and use committees (IACUC). MUC16-mediated tumor growth: T3M4 WT, WT-MUC16^{KO} (2E4), T3M4 SC, and SC-MUC16^{KO} (1E10) cells ($3 \times 10^5/30$ μ L PBS) were orthotopically implanted into the pancreas of athymic nu/nu mice (Crl:NU-Foxn1nu; n = 14 each group) and the tumor-bearing animals were sacrificed (day 28) and analyzed as described previously.²⁷ mAb AR9.6 treatment: to determine the early preclinical response of AR9.6 in orthotopic tumor model, T3M4 WT and SC cells ($2.5\text{--}3 \times 10^5/30$ μ L PBS) implanted tumor-bearing athymic nu/nu mice were randomized (after 14 days) and treated with four doses of vehicle control (PBS, n = 11) and AR9.6 mAb (0.5 mg, n = 13) via intraperitoneal injection (i.p.) with 72 h interval for four cycles. mAb AR9.6 plus GEM combination treatment: to determine the effect of combination therapy of mAb AR9.6 with GEM at later stages of tumor growth, T3M4 WT cells ($3 \times 10^5/30$ μ L PBS) implanted tumor-bearing athymic nu/nu mice were randomized (after 18 days) and treated as follows: group I, vehicle control (PBS, n = 9); group II, IgG (0.5 mg, n = 9); group III, GEM (60 mg/kg, n = 10); group IV, mAb AR9.6 (0.5 mg, n = 10), and group V, mAb AR9.6 (0.5 mg) + GEM (60 mg/kg, n = 10) four doses via i.p. with 72 h interval for four cycles. At the end of each study, the animals were sacrificed, and the tumor weight, volume, and incidence of metastases were determined.

Western blotting

T3M4 (WT, WT-MUC16^{KO}, SC, SC-MUC16^{KO}) and Capan-2 (WT, WT-MUC16^{KO}, SC, SC-MUC16^{KO}) cell lysates (radioimmunoprecipitation assay [RIPA] lysis buffer [Thermo Scientific, Rockford, IL USA] with protease and proteinase inhibitor) were resolved on 4%–20% SDS-PAGE (30–50 μ g/lane) and transferred to polyvinylidene fluoride (PVDF) membrane (Thermo Scientific, Rockford, IL, USA). For analyzing the effect of various anti-MUC16 antibodies, T3M4 WT and SC cells were treated with IgG, 5E11 (provided by Dr. Ulla Mandel, University of Copenhagen, Denmark), B43.13 and AR9.6 (2.5 and 5 μ g/mL; Quest PharmaTech, Canada) for 24 h. For analyzing the effect of TR1.2, T3M4 SC-MUC16^{KO} cells (polyclonal) were treated with TR1.2 (1 and 2 μ g/mL) for 14 h. For analyzing the effect of TR1.2 and AR9.6, T3M4 WT-MUC16^{KO} cells were treated with TR1.2 (1 μ g/mL) with or without mAb AR9.6 (5 μ g/mL) for 14 h. At the end of each experimental time point, the cell lysates were prepared, and proteins were separated as mentioned above. The membranes were probed with the following primary and secondary antibodies as per the manufacturer's recommendations: anti-p-AKT (S473), anti-AKT, anti-p-GSK3 β (S9), anti-GSK3 β , anti-p-ErbB2 (Y1248), anti-ErbB2, anti-p-ErbB3 (Y1289), anti-ErbB3, anti-cyclin D1, anti-cyclin E1, anti-GAPDH, anti- β -actin (Cell Signaling Technology, Danvers,

MA, USA), anti-p-ErbB1 (Y1173; Santa Cruz Biotechnology, CA, USA), and horseradish peroxidase (HRP) conjugated goat anti-mouse IgG and goat anti-rabbit IgG (Cell Signaling Technology, Danvers, MA, USA). The signal was detected by using the enhanced chemiluminescent kit (Bio-Rad, Hercules, CA, USA).

IHC

The standard IHC method was used for the analysis of protein expression in tissue samples. Briefly, the paraffin-embedded slides were deparaffinized with xylene, re-hydrated with series of alcohol (100% to 70%), quenched with hydrogen peroxide, and immersed in 95°C citrate buffer (pH = 6.0) for 15 min for antigen retrieval. After blocking with a universal blocker (Thermo Fisher Scientific, Waltham, MA, USA), the slides were incubated with antibodies OC125 (1:100; Abcam, Cambridge, MA, USA) AR9.6 (1:500), 5E11 (1 µg/mL), and 5B9 (1 µg/mL; provided by Dr. Ulla Mandel, University of Copenhagen, Denmark). The slides were washed and incubated with HRP conjugated secondary antibody. After 1 h, the slides were washed with Tris-buffered saline (TBS), and the antigen-antibody reaction products were developed using 3,3'-diaminobenzidine tetrahydrochloride (DAB, Vector Laboratories, Burlingame, CA, USA) substrate and then counterstained with hematoxylin. All the slides were dehydrated with alcohol series, and after xylene washes, the slides were mounted with the coverslip. Also, analysis of MUC16 expression in a pancreatic cancer tissue microarray (BIC14011a; BIOMAX.US) was performed as described above. For the analysis of p-ErbB2 (1:50; Abcam, Cambridge, MA, USA) and p-AKT (1:50; Abcam, Cambridge, MA, USA) in mouse tissue sections, the slides were processed as mentioned above. The protein expression was scored by a pathologist. The histological scoring was performed based on stain proportion (0%–100%) and intensity (0, negligible; 1, low; 2, moderate; 3, high). The histoscore was generated by multiplying the stain proportion score (1, <5%; 2, 5%–25%; 3, 26%–50%; 4, 51%–75%; 5, >75%) with the intensity score (0–3) to obtain values between 0 and 15.

Proliferation index

Tissue sections of T3M4 cells implanted tumors treated with vehicle control (PBS), IgG, GEM, mAb AR9.6, and mAb AR9.6 + GEM (n = 9) were stained with rabbit anti-Ki-67 antibody (Abcam, Cambridge, MA, USA) as described previously.²⁸ A total of four random fields were examined in nine different tumor samples from each group. The results were expressed as a mean percentage of Ki-67-positive cells ± SEM per field (100× magnification).

Microvessel density

Tissue sections of T3M4 cells implanted tumors treated with vehicle control (PBS), IgG, GEM, mAb AR9.6, and mAb AR9.6 + GEM (n = 9) were stained with rabbit anti-CD31 antibody (Abcam, Cambridge, MA, USA), as described previously.²⁸ The number of the CD31-positive vessel was examined under a microscope and counted. A total of four random fields were examined in nine different tumor samples from each group. The results were expressed as a mean percentage of vessels ± SEM per field (100× magnification).

Proximity ligation assay

The protein-protein interaction experiments (growth factor receptors interactions with mucins and integrin complexes) were performed by using Duo Link II (Olink Bioscience, Watertown, MA, USA) Proximity Ligation Assay kit according to manufacturer's recommendation as described previously.²⁹ Briefly, T3M4 WT and SC cells were grown on coverslips and then incubated with the primary antibody specific to MUC16 (AR9.6, Quest PharmaTech, Canada) and ErbB2 (Santa Cruz Biotechnology, CA, USA) and then the primary antibodies were incubated with PLA probes anti-Rabbit PLUS, anti-mouse MINUS. Experiments with omitting the primary antibody used for single recognition PLA or either one of the primary antibodies used in double recognition PLA served as a negative control. The fluorescence images were captured under Zeiss LSM 510 laser scanning confocal microscope (Carl Zeiss, Thornwood, NY, USA) at Confocal Laser Scanning Fluorescence Microscopy Core Facility, UNMC, and the signal/dots per cells were quantified by using blob finder software.

Epitope mapping

The recombinant MUC16 constructs (TR1.7, TR1.2, 1/4, 5/2, 5/7, 5/8, 5/9, K292, and F/R) were synthesized and expressed, and ELISA assays were performed as described previously.¹ Briefly, MUC16 recombinant proteins were serially diluted from an initial concentration of 500 ng/mL and coated overnight at 4°C in carbonate-bicarbonate buffer (pH 9.6) in a Nunc MaxiSorp 96-well plates (Thermo Scientific, Rockford, IL USA). Then the plate was blocked with PLI-P buffer pH 7.4 (PO₄, Na/K, 1% Triton, 1% BSA) for 1 h at room temperature, and incubated with mAb AR9.6 (Quest PharmaTech, Canada) at 1 µg/mL for 1 h at room temperature. Bound antibodies were detected with HRP-conjugated IgG goat anti-mouse antibody (Southern Biotech, Birmingham, AL, USA). Plates were developed with 3,3',5,5'-Tetramethylbenzidine (TMB⁺) substrate (Dako, Carpinteria, CA, USA), reactions stopped with 0.5 M H₂SO₄, and absorbance read at 450 nm. Incubation of plates with anti-MUC16 mAb 5E11 served as a positive control, and plates incubated with anti-GalNAc-T3 mAb 2D10 (IgG) were used as a negative control.

Live/dead cytotoxicity assay

T3M4 and Capan-2 cells were treated with vehicle (PBS), IgG (10 µg/mL), AR9.6 (10 µg/mL), GEM (15 nM), and AR9.6, + GEM for 24 h and have performed live/dead cytotoxicity assay as per the manufacturer's instruction (Invitrogen, CA, USA). Briefly, after washing with PBS, the cells were incubated with assay reagent containing 20 µL ethidium homodimer-1 (EthD-1, 2 mM) and 5 µL of calcein acetoxymethyl ester (Calcein-AM, 4 mM) in 10 mL PBS for 30 min. Images of live (green) and dead (red) cells were captured with Nikon Eclipse Ts2R fluorescent microscope and counted with ImageJ software (n = 3).

Statistical analysis

For unmatched samples, the continuous characteristics and outcomes were compared using t tests or nonparametric Wilcoxon rank-sum tests between two groups, using two-way ANOVA with Tukey's multiple comparisons or nonparametric Kruskal-Wallis tests with

Bonferroni's multiple comparisons between more than two groups; the categorical characteristics and outcomes were compared using chi-square tests or Fisher's exact tests between groups, and Bonferroni's correction was used for multiple comparisons if needed. For matched samples, the continuous characteristics and outcomes were compared using paired t tests or nonparametric Wilcoxon signed-rank tests between two groups. The cell proliferation was compared between the groups by two-way ANOVA with Tukey's multiple comparisons at each time points. Tumor cell migration, invasion, *in vivo* tumor weight, tumor volume, Ki-67, CD31, and live/dead cell assay was analyzed by one-way ANOVA with Dunnett's multiple comparison test. Patient survival was plotted using the method of Kaplan-Meier and compared between groups using log-rank (Mantel-Cox) tests. Mann-Whitney rank-sum test was used between the PDAC subtype classification system. A value of $p < 0.05$ was considered to be statistically significant.

SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at <https://doi.org/10.1016/j.ymthe.2020.12.029>.

ACKNOWLEDGMENTS

This work was supported by the National Cancer Institute at the National Institutes of Health R01 CA208108, NIH/SPORE Carrier Development Award (P50CA127297), and NE-DHHS/LB506 to P.R.; NIH/SPORE P50CA127297, Program Project Grant P01CA217798, Early Detection Research Network 5U01CA111294, Pancreatic Cancer Detection Consortium 5U01CA210240, and Tumor Microenvironment Network U54 CA163120 to M.A.H. and R50CA211462 to P.M.G.; The National Cancer Institute at the National Institutes of Health R01CA199064 to J.J.Y.; and The Danish National Research Foundation DNR107 to H.C. Also, we thank Jessica Odvody, Premila Leiphakpam, Krishti Sabloak, and Philamon Leon for their technical assistance.

AUTHOR CONTRIBUTIONS

P.R. conceived and supervised the study. D.T., S.S., and X.L. performed most of the *in vitro* and *in vivo* experiments. H.-R.L., J.A.G., P.M.G., and T.C. performed orthotopic tumor studies. B.S. scored the immunohistochemical staining. L.M.-S., H.C., C.S., and H.H.W. provided COSMC KO PDAC cells and performed anti-MUC16 mAb epitope mapping studies. H.C.M. and K.P.O. performed RNA sequencing analysis. X.L.P. and J.J.Y. performed tumor sub-type analysis with MUC16 expression. F.Q. and F.Y. performed biostatistical analysis. R.M. provided AR9.6 mAb expressing clones. U.M. provided anti-MUC16 antibodies (5E11 and 5B9) for the study. M.A.H. analyzed the data. D.T., M.A.H., and P.R. wrote the manuscript.

DECLARATION OF INTERESTS

M.A.H. and P.R. have an equity interest in OncoCare Therapeutics. R.M. is employed by Quest PharmaTech and has an equity interest in this company. All other authors declare no competing interests.

REFERENCES

- Marcos-Silva, L., Narimatsu, Y., Halim, A., Campos, D., Yang, Z., Tarp, M.A., Pereira, P.J.B., Mandel, U., Bennett, E.P., Vakhrushev, S.Y., et al. (2014). Characterization of binding epitopes of CA125 monoclonal antibodies. *J. Proteome Res.* *13*, 3349–3359.
- Remmers, N., Anderson, J.M., Linde, E.M., DiMaio, D.J., Lazenby, A.J., Wandall, H.H., Mandel, U., Clausen, H., Yu, F., and Hollingsworth, M.A. (2013). Aberrant expression of mucin core proteins and o-linked glycans associated with progression of pancreatic cancer. *Clin. Cancer Res.* *19*, 1981–1993.
- Liang, C., Qin, Y., Zhang, B., Ji, S., Shi, S., Xu, W., Liu, J., Xiang, J., Liang, D., Hu, Q., et al. (2017). Oncogenic KRAS Targets MUC16/CA125 in Pancreatic Ductal Adenocarcinoma. *Mol. Cancer Res.* *15*, 201–212.
- O'Brien, T.J., Beard, J.B., Underwood, L.J., Dennis, R.A., Santin, A.D., and York, L. (2001). The CA 125 gene: an extracellular superstructure dominated by repeat sequences. *Tumour Biol.* *22*, 348–366.
- Hakomori, S. (2002). Glycosylation defining cancer malignancy: new wine in an old bottle. *Proc. Natl. Acad. Sci. USA* *99*, 10231–10233.
- Pinho, S.S., and Reis, C.A. (2015). Glycosylation in cancer: mechanisms and clinical implications. *Nat. Rev. Cancer* *15*, 540–555.
- Julien, S., Videira, P.A., and Delannoy, P. (2012). Sialyl-Tn in cancer: (how) did we miss the target? *Biomolecules* *2*, 435–466.
- Kim, G.E., Bae, H.I., Park, H.U., Kuan, S.F., Crawley, S.C., Ho, J.J.L., and Kim, Y.S. (2002). Aberrant expression of MUC5AC and MUC6 gastric mucins and sialyl Tn antigen in intraepithelial neoplasms of the pancreas. *Gastroenterology* *123*, 1052–1060.
- Radhakrishnan, P., Dabelsteen, S., Madsen, F.B., Francavilla, C., Kopp, K.L., Steentoft, C., Vakhrushev, S.Y., Olsen, J.V., Hansen, L., Bennett, E.P., et al. (2014). Immature truncated O-glycophenotype of cancer directly induces oncogenic features. *Proc. Natl. Acad. Sci. USA* *111*, E4066–E4075.
- Steentoft, C., Vakhrushev, S.Y., Joshi, H.J., Kong, Y., Vester-Christensen, M.B., Schjoldager, K.T.-B.G., Lavrsen, K., Dabelsteen, S., Pedersen, N.B., Marcos-Silva, L., et al. (2013). Precision mapping of the human O-GalNAc glycoproteome through SimpleCell technology. *EMBO J.* *32*, 1478–1488.
- Haridas, D., Chakraborty, S., Ponnusamy, M.P., Lakshmanan, I., Rachagani, S., Cruz, E., Kumar, S., Das, S., Lele, S.M., Anderson, J.M., et al. (2011). Pathobiological implications of MUC16 expression in pancreatic cancer. *PLoS ONE* *6*, e26839.
- Law, C.W., Chen, Y., Shi, W., and Smyth, G.K. (2014). voom: Precision weights unlock linear model analysis tools for RNA-seq read counts. *Genome Biol.* *15*, R29.
- Marcos-Silva, L., Ricardo, S., Chen, K., Blixt, O., Arigi, E., Pereira, D., Høgdall, E., Mandel, U., Bennett, E.P., Vakhrushev, S.Y., et al. (2015). A novel monoclonal antibody to a defined peptide epitope in MUC16. *Glycobiology* *25*, 1172–1182.
- Cancer Genome Atlas Research Network. Electronic address: andrew_aguirre@dfci.harvard.edu; Cancer Genome Atlas Research Network (2017). Integrated Genomic Characterization of Pancreatic Ductal Adenocarcinoma. *Cancer Cell* *32*, 185–203.e13.
- Noujaim, A.A., Schultes, B.C., Baum, R.P., and Madiyalakan, R. (2001). Induction of CA125-specific B and T cell responses in patients injected with MAb-B43.13—evidence for antibody-mediated antigen-processing and presentation of CA125 *in vivo*. *Cancer Biother. Radiopharm.* *16*, 187–203.
- Balachandran, V.P., Luksza, M., Zhao, J.N., Makarov, V., Moral, J.A., Remark, R., Herbst, B., Askan, G., Bhanot, U., Senbabaoglu, Y., et al.; Australian Pancreatic Cancer Genome Initiative; Garvan Institute of Medical Research; Prince of Wales Hospital; Royal North Shore Hospital; University of Glasgow; St Vincent's Hospital; QIMR Berghofer Medical Research Institute; University of Melbourne; Centre for Cancer Research; University of Queensland, Institute for Molecular Bioscience; Bankstown Hospital; Liverpool Hospital; Royal Prince Alfred Hospital; Chris O'Brien Lifehouse; Westmead Hospital; Fremantle Hospital; St John of God Healthcare; Royal Adelaide Hospital; Flinders Medical Centre; Envoi Pathology; Princess Alexandria Hospital; Austin Hospital; Johns Hopkins Medical Institutes; ARC-Net Centre for Applied Research on Cancer (2017). Identification of unique neoantigen qualities in long-term survivors of pancreatic cancer. *Nature* *551*, 512–516.

17. Chen, S.H., Hung, W.C., Wang, P., Paul, C., and Konstantopoulos, K. (2013). Mesothelin binding to CA125/MUC16 promotes pancreatic cancer cell motility and invasion via MMP-7 activation. *Sci. Rep.* 3, 1870.
18. Normanno, N., De Luca, A., Bianco, C., Strizzi, L., Mancino, M., Maiello, M.R., Carotenuto, A., De Feo, G., Caponigro, F., and Salomon, D.S. (2006). Epidermal growth factor receptor (EGFR) signaling in cancer. *Gene* 366, 2–16.
19. Bailey, P., Chang, D.K., Nones, K., Johns, A.L., Patch, A.M., Gingras, M.C., Miller, D.K., Christ, A.N., Bruxner, T.J.C., Quinn, M.C., et al.; Australian Pancreatic Cancer Genome Initiative (2016). Genomic analyses identify molecular subtypes of pancreatic cancer. *Nature* 531, 47–52.
20. Eser, S., Reiff, N., Messer, M., Seidler, B., Gottschalk, K., Dobler, M., Hieber, M., Arbeiter, A., Klein, S., Kong, B., et al. (2013). Selective requirement of PI3K/PDK1 signaling for Kras oncogene-driven pancreatic cell plasticity and cancer. *Cancer Cell* 23, 406–420.
21. Schlieman, M.G., Fahy, B.N., Ramsamooj, R., Beckett, L., and Bold, R.J. (2003). Incidence, mechanism and prognostic value of activated AKT in pancreas cancer. *Br. J. Cancer* 89, 2110–2115.
22. Yamamoto, S., Tomita, Y., Hoshida, Y., Morooka, T., Nagano, H., Dono, K., Umeshita, K., Sakon, M., Ishikawa, O., Ohigashi, H., et al. (2004). Prognostic significance of activated Akt expression in pancreatic ductal adenocarcinoma. *Clin. Cancer Res.* 10, 2846–2850.
23. Hollingsworth, M.A., and Swanson, B.J. (2004). Mucins in cancer: protection and control of the cell surface. *Nat. Rev. Cancer* 4, 45–60.
24. Schjoldager, K.T., and Clausen, H. (2012). Site-specific protein O-glycosylation modulates proprotein processing - deciphering specific functions of the large polypeptide GalNAc-transferase gene family. *Biochim. Biophys. Acta* 1820, 2079–2094.
25. Steentoft, C., Vakhrushev, S.Y., Vester-Christensen, M.B., Schjoldager, K.T.-B.G., Kong, Y., Bennett, E.P., Mandel, U., Wandall, H., Lavery, S.B., and Clausen, H. (2011). Mining the O-glycoproteome using zinc-finger nuclease-glycoengineered SimpleCell lines. *Nat. Methods* 8, 977–982.
26. Radhakrishnan, P., Grandgenett, P.M., Mohr, A.M., Bunt, S.K., Yu, F., Chowdhury, S., and Hollingsworth, M.A. (2013). Expression of core 3 synthase in human pancreatic cancer cells suppresses tumor growth and metastasis. *Int. J. Cancer* 133, 2824–2833.
27. Tsutsumida, H., Swanson, B.J., Singh, P.K., Caffrey, T.C., Kitajima, S., Goto, M., Yonezawa, S., and Hollingsworth, M.A. (2006). RNA interference suppression of MUC1 reduces the growth rate and metastatic phenotype of human pancreatic cancer cells. *Clin. Cancer Res.* 12, 2976–2987.
28. Radhakrishnan, P., Bryant, V.C., Blowers, E.C., Rajule, R.N., Gautam, N., Anwar, M.M., Mohr, A.M., Grandgenett, P.M., Bunt, S.K., Arnst, J.L., et al. (2013). Targeting the NF- κ B and mTOR pathways with a quinoxaline urea analog that inhibits IKK β for pancreas cancer therapy. *Clin. Cancer Res.* 19, 2025–2035.
29. Söderberg, O., Gullberg, M., Jarvius, M., Ridderstråle, K., Leuchowius, K.J., Jarvius, J., Wester, K., Hydbring, P., Bahram, F., Larsson, L.G., and Landegren, U. (2006). Direct observation of individual endogenous protein complexes in situ by proximity ligation. *Nat. Methods* 3, 995–1000.

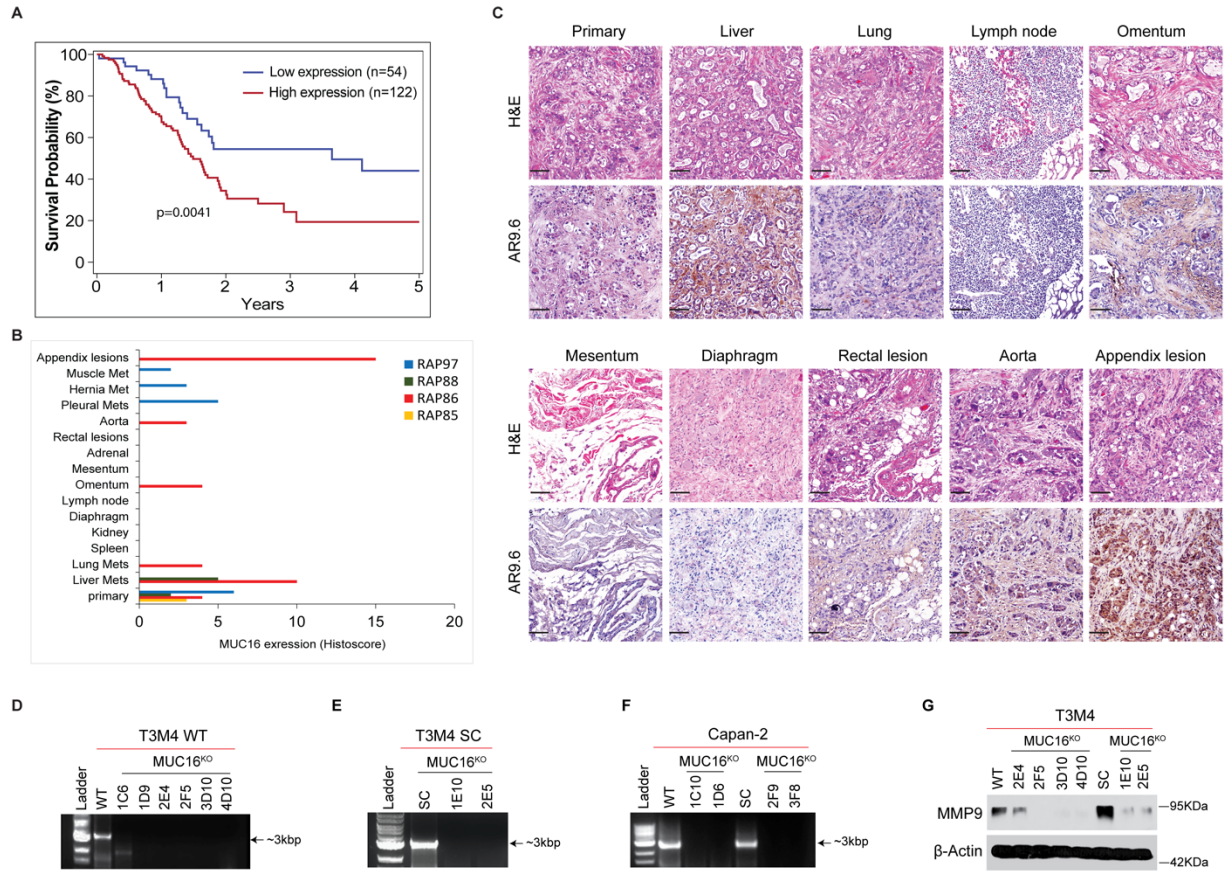
Supplemental Information

Isoforms of MUC16 activate oncogenic signaling through EGF receptors to enhance the progression of pancreatic cancer

Divya Thomas, Satish Sagar, Xiang Liu, Hye-Rim Lee, James A. Grunkemeyer, Paul M. Grandgenett, Thomas Caffrey, Kelly A. O'Connell, Benjamin Swanson, Lara Marcos-Silva, Catharina Steentoft, Hans H. Wandall, Hans Carlo Maurer, Xianlu Laura Peng, Jen Jen Yeh, Fang Qiu, Fang Yu, Ragupathy Madiyalakan, Kenneth P. Olive, Ulla Mandel, Henrik Clausen, Michael A. Hollingsworth, and Prakash Radhakrishnan

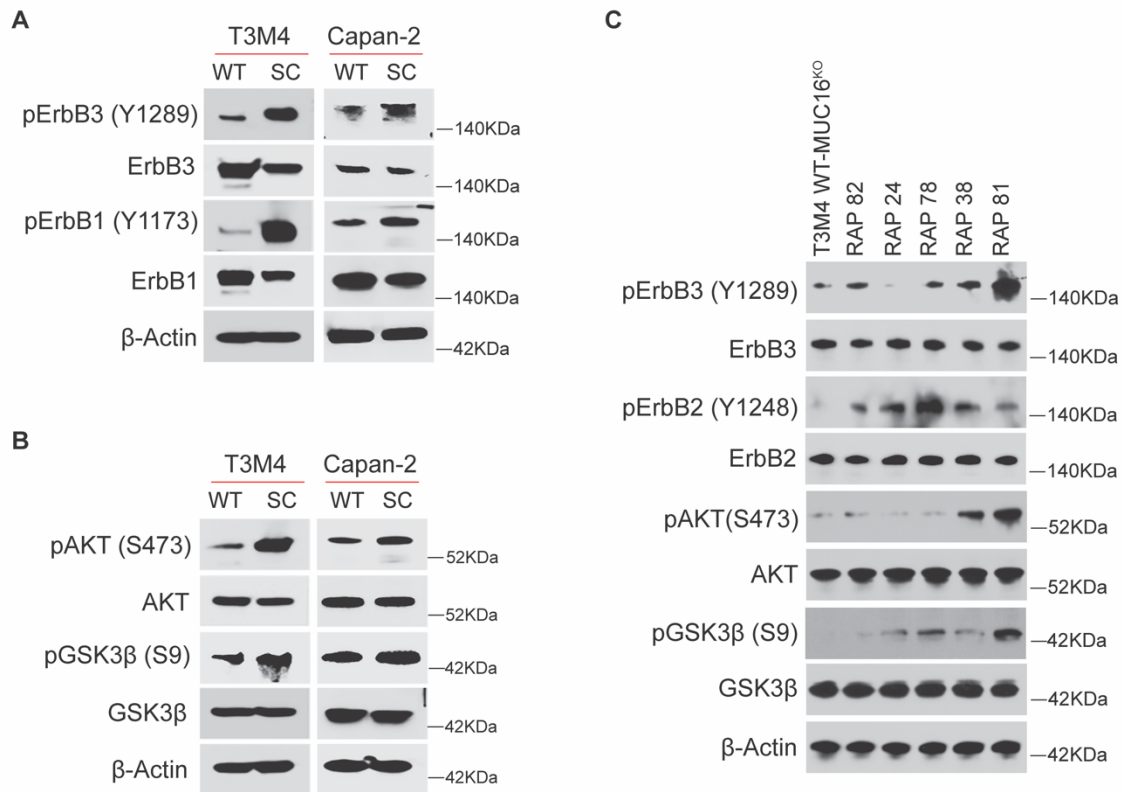
Supplemental Figures

Supplementary Figure 1



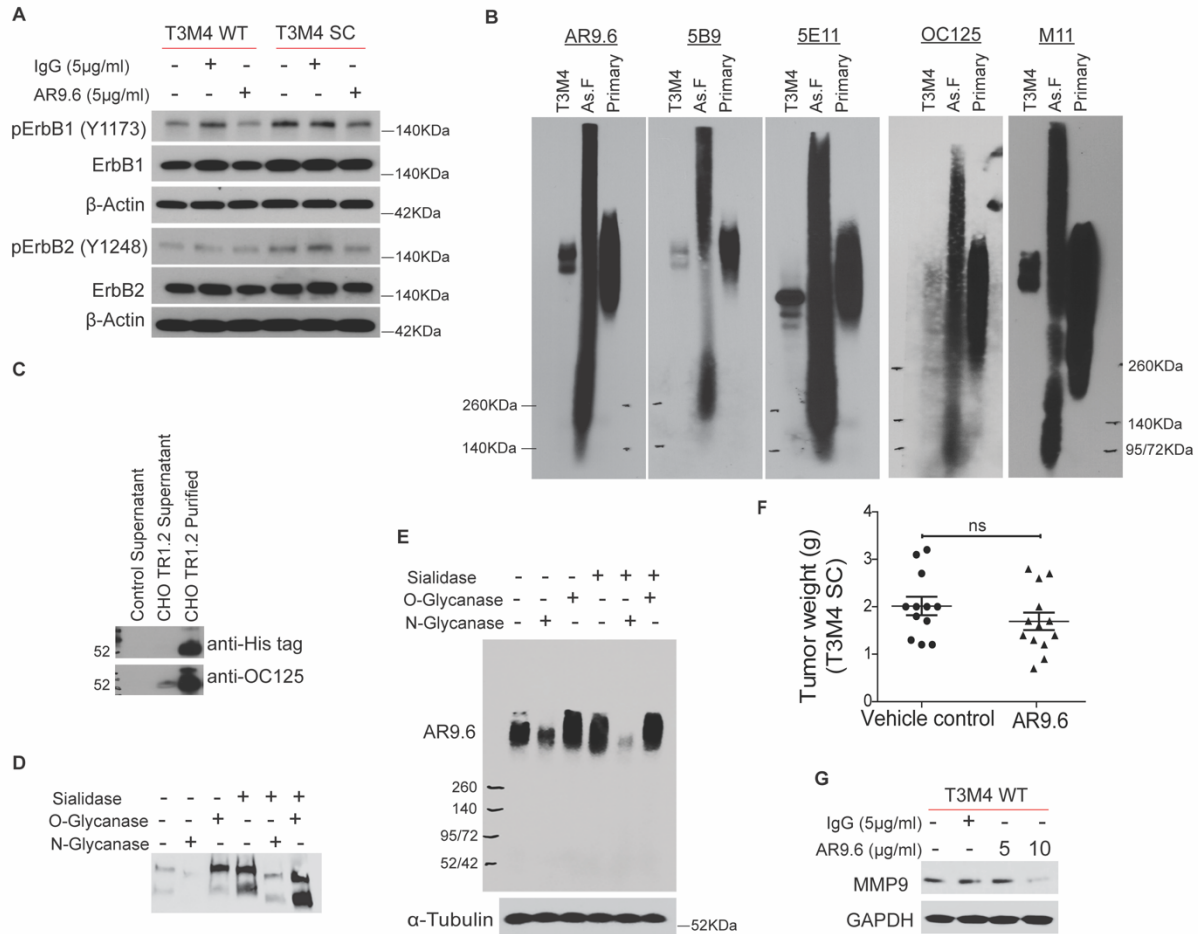
Supplemental Figure S1. MUC16 expression in PDAC. **A.** MUC16 mRNA expression and PDAC patient survival probability was analyzed from the Human Protein Atlas. Kaplan-Meier plot evaluated survival probability of patients expressing low MUC16 (n=54) and high MUC16 (n=122). A significant difference was observed between the two groups at $p=0.0041$ (log-rank test). **B.** Histscore of IHC analysis of MUC16 expression in selected RAP patients with multiple metastatic sites (n=4) using MUC16 specific mAb AR 9.6. **C.** Representative images of IHC analysis of MUC16 expression in selected RAP patients with multiple metastatic sites using MUC16 specific mAb AR 9.6. Scale bar = 40 μ m. **Genetic deletion of MUC16 in PDAC cells. D-F.** CRISPR/Cas9 mediated genetic deletion of MUC16 in T3M4 and Capan-2 (WT and SC) cells and detected by genomic DNA PCR. Parental cells show amplification of a band at 3000bp, which was deleted in MUC16^{KO} clones of PDAC cells. **G.** Western blotting analysis of MMP9 in T3M4 WT, WT-MUC16^{KO}, SC and SC-MUC16^{KO} cells. Detection of β -actin served as the loading control.

Supplementary Figure 2



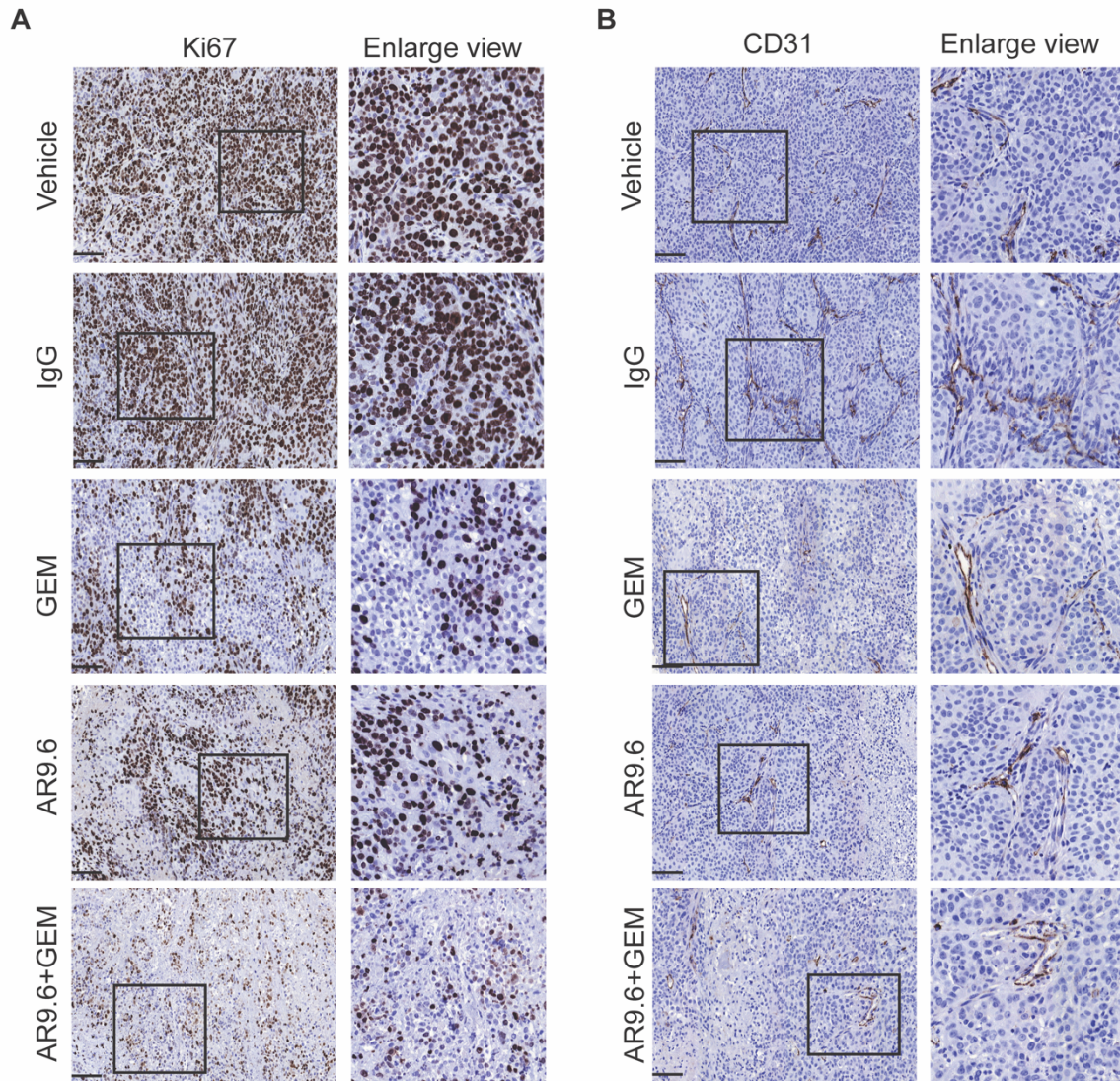
Supplemental Figure S2. Interaction of MUC16 with ErbB receptors. **A.** Western blotting analysis of p-ErbB3 (Y1289), ErbB3, p-ErbB1 (Y1173) and ErbB1 in T3M4 and Capan-2 WT and SC cells. **B.** Western blotting analysis of p-AKT (S473), AKT, p-GSK3β (S9) and GSK3β in T3M4 and Capan2 WT and SC cells. **C.** Western blotting analysis of p-ErbB3 (Y1289), ErbB3, p-ErbB2 (Y1248) and ErbB2, p-AKT (S473), AKT, p-GSK3β (S9) and GSK3β in T3M4WT-MUC16^{KO} cells treated with purified MUC16 (5 μg/ml for 24 h) from PDAC patients (n=5). Detection of β-actin served as the loading control.

Supplementary Figure 3



Supplemental Figure S3. Characterization of MUC16 TR1.2 epitope. **A.** Western blotting of p-ErbB1 (Y1173), ErbB1, p-ErbB2 (Y1248) and ErbB2 in mAb AR9.6 treated T3M4 WT and SC cells. **B.** SDS-agarose western blotting analysis of MUC16 in T3M4 cells lysate, ascetic fluid (As.F) (RAP # 81) and pancreas tumor tissue lysates (RAP # 35) from the PDAC patients samples using MUC16 specific antibodies mAb AR9.6, 5B9, 5E11, OC125 and M11. **C.** MUC16 TR1.2 purified from CHO cells and probed with anti-His-tag and OC125 antibodies. Treatment of MUC16 TR1.2 glycopeptides (**D**) and T3M4 cell lysates (**E**) with Sialidase, O-Glycanase and N-Glycanase, and probed with AR9.6 mAb. Detection of α -tubulin served as the loading control. **F.** T3M4 SC cells implanted tumor-bearing animals were treated with Vehicle control (n=12) and mAb AR9.6 (n=13). [ns, not significant]. **G.** Western blotting analysis of MMP9 in vehicle, mouse IgG and mAb AR9.6 treated T3M4 cells. Detection of β -actin and GAPDH served as the loading control.

Supplementary Figure 4



Supplemental Figure S4. mAb AR9.6 plus Gemcitabine treatment reduces *in vivo* tumor growth. Representative images of immunohistochemical analysis of Ki-67 (**A**) and CD31 (**B**) in T3M4 tumor-bearing mice treated with PBS, IgG, gemcitabine, mAb AR9.6, and mAb AR9.6 plus Gemcitabine (n=9).

Supplemental Tables

Table S1. Clinical information of UNMC Rapid Autopsy Pancreas Cancer Patients

RAP #	Age	Sex	Survival days
03	78	F	274
04	59	M	110
05	65	F	35
06	62	M	549
07	71	M	225
08	72	M	9
09	69	M	198
10	74	M	85
11	80	M	185
12	82	M	364
13	72	M	42
16	70	M	178
19	65	M	145
20	77	M	309
21	60	F	221
25	74	M	206
29	80	F	100
30	55	M	148
31	79	M	21
32	59	M	288
33	50	M	228
35	62	M	419
37	58	M	293
39	75	M	65
40	62	M	336
44	58	F	386
46	78	F	71
50	47	F	276
51	57	M	1422
53	57	M	443
54	68	M	168
55	55	F	669
56	74	M	364
57	56	M	51
58	60	M	646
59	62	F	495
61	78	M	269
62	75	M	203

63	70	M	865
64	55	M	423
66	56	M	247
68	66	M	635
72	78	M	366
73	52	M	191
77	80	M	394
78	63	M	451
81	60	F	182
82	71	M	595
85	67	M	446
86	43	M	64
87	56	M	409
88	67	M	227
91	54	M	2011
94	80	M	2282
97	67	M	217
99	70	M	966
100	61	M	1228
101	64	F	725
104	53	M	263
105	89	F	26
109	64	M	405
119	85	F	183
120	69	M	507
123	74	M	90
125	49	F	27
127	81	M	150
129	75	F	427
130	70	M	15
131	72	M	1360
132	77	F	75
133	64	F	1176
135	84	M	893

Table S2. Statistical comparison of IHC histoscores between different anti-MUC16 antibodies.

Trial	N Obs	N	Mean	Std Dev	Median	Minimum	Maximum	P value when Comparing with OC125	Overall P value
OC125	61	61	0.84	1.00	0	0	3.00	Ref	0.002
AR9.6	61	61	1.34	1.18	1	0	3.00	<0.0001	
5B9	61	61	1.15	1.03	1	0	3.00	0.016	
5E11	61	61	0.92	1.00	1	0	3.00	0.40	

UNMC RAP PDAC patients samples. The intensity data were summarized using median value since the data does not follow normal distribution. The Friedman test showed significant differences in the median intensities among at least two of the four antibodies in the studied samples ($p=0.002$). Specifically, the antibody 5E9 (p value=0.016) and AR9.6 (p value<0.0001) shows significantly higher values than OC125.

Table S3. PDAC subtype classification

Classification	Subtype	MUC16
Bailey	Squamous vs Immunogenic	0.00057506
	Squamous vs Progenitor	5.31E-05
	Squamous vs ADEX	8.97E-05
	Immunogenic vs Progenitor	4.24E-01
	Immunogenic vs ADEX	6.66E-01
	Progenitor vs ADEX	7.51E-01
Collisson	Classical vs Exocrine-like	0.8659441
	Classical vs Quasimesenchymal	0.09727453
	Exocrine-like vs Quasimesenchymal	0.05695079
Moffit	Basal-like vs Classical	2.89E-11

Mann-Whitney rank-sum test was used between the PDAC subtype classification system. A p value of less than 0.01 was considered statistically significant.

Table S4. The incidence of tumor metastasis.

T3M4 Cells	Peritoneum	Diaphragm	Lymph node	Liver	Lung	Invaded spleen
WT	35.71% (5/14)	28.57% (4/14)	21.42% (3/14)	42.85% (6/14)	21.42% (3/14)	85.71% (12/14)
WT-MUC16 ^{KO}	21.42% (3/14)	7.1% (1/14)	14.28% (2/14)	7.1% (1/14)	0% (0/14)	42.85% (6/14)
SC	64.28% (9/14)	50% (7/14)	57.14% (8/14)	57.14% (8/14)	42.85% (6/14)	85.71% (12/14)
SC-MUC16 ^{KO}	14.28% (2/14)	7.1% (1/14)	14.28% (2/14)	0% (0/14)	0% (0/14)	28.57% (4/14)
WT vs WT-MUC16^{KO}	<i>0.99</i>	<i>0.98</i>	<i>0.99</i>	<i>0.23</i>	<i>0.66</i>	<i>0.054</i>
WT vs SC	<i>0.39</i>	<i>0.74</i>	<i>0.16</i>	<i>0.99</i>	<i>0.99</i>	<i>0.99</i>
SC vs SC-MUC16^{KO}	<i>0.02</i>	<i>0.099</i>	<i>0.054</i>	<i>0.006</i>	<i>0.048</i>	<i>0.007</i>

Chi-square test or Fisher's exact test was used to compare incidence of tumor metastasis between the following groups: WT vs WT-MUC16^{KO}, WT vs SC, and SC vs SC-MUC16^{KO}. Bonferroni's correction was used for multiple comparisons. A p-value of < 0.05 was considered statistically significant

Table S5. Ingenuity Pathway Analysis (IPA).

Top Canonical Pathways		
Name	p-value	Ratio
Molecular Mechanisms of Cancer	1.4E-16	12/381 (0.031)
B Cell Receptor Signaling	3.04E-16	10/171 (0.058)
ErbB2-ErbB3 Signaling	6.56E-16	8/60 (0.133)
14-3-3-mediated Signaling	2.05E-15	9/121 (0.074)
Prostate Cancer Signaling	1.54E-14	8/99 (0.081)

Previous global profiling of phosphoprotein expression patterns and signal transduction pathways of cells with enforced deletion of COSMC revealed a constitutive activation of steady-state oncogenic signaling cascades¹. Examination of those data by IPA and deductive reasoning identified ErbB2-ErbB3 and FAK as principal nodes for many of the observed effects of truncated O-glycosylation on tumor malignancy.

Supplemental Methods

PDAC Patients Survival Analysis

MUC16 mRNA expression and PDAC patient's survival was analyzed from Human Protein Atlas: <https://www.proteinatlas.org/ENSG00000181143-MUC16/pathology/tissue/pancreatic+cancer> based upon the data generated by the TCGA Research Network. Kaplan plot and the log-rank *p-value* was generated with patients with low MUC16 expression (n=54) and patients with high MUC16 expression (n=122).

LCM-RNA sequencing of Epithelial and Stromal PDAC tissues

Sample acquisition

Freshly frozen tissue samples of PDAC were obtained from surgical specimens from patients undergoing surgery at Pancreas Center at Columbia University Medical Center. Prior to surgery, all patients provided informed consent, which was approved by the local Institutional Review Board (IRB protocol #AAAB2667). Immediately after surgical removal, the specimens were sectioned and microscopically evaluated by the Columbia University Tumor Bank. Suitable samples were transferred into optimal cutting temperature (OCT) medium (Tissue Tek) and snap frozen in a 2-methylbutane dry ice slurry. The tissue blocks were stored at -80°C until further processing. PDAC samples in the Tumor Bank were screened for diagnosis, purity, and viability by H&E analysis of frozen blocks. The overall sample RNA quality was initially assessed by gel electrophoresis; samples exhibiting high RNA quality were utilized for subsequent analyses.

Sample extraction

Frozen tissue specimens were cut at 8 to 9µm thickness and 2 to 3 sections were transferred onto a polyethylene naphthalate membrane glass slide (Arcturus, Applied Biosystems). For initial histopathological review, immediately adjacent sections were cut and stained using a standard Hematoxylin and Eosin protocol to confirm the diagnosis and selected regions of interest.

Laser capture microdissection

Throughout the staining procedure, RNase-free water was used. Fixation was done with 95% ethanol, followed by Crystal violet acetate staining (1% in Tris-buffered 70% ethanol), a brief washing step in 70% ethanol and final dehydration in 100% ethanol. Laser capture microdissection was performed on a PALM MicroBeam microscope (Zeiss) to collect at least 1000 cells per compartment from selected (pre-)malignant areas.

RNA

RNA was extracted using the RNeasy Plus Micro Kit (Qiagen, USA) following the manufacturer's instructions. Prior to processing further, RNA integrity and yield were determined using the Agilent 2100 Bioanalyzer (RNA 6000 Pico Kit for LCM and RNA 6000 Nano Kit for bulk samples, respectively). Yields ranged from 1 to 10 ng per LCM sample and several µg per bulk sample, respectively. Only samples with an RNA Integrity Number (RIN) of at least 7 were used for further processing.

RNA amplification and library preparation

1 - 2 ng of RNA from LCM samples were amplified using the Ovation RNA-Seq System V2 Kit (NuGEN, CA, USA) following the manufacturer's instructions. The resulting cDNA libraries were fragmented using a Covaris S2 Sonicator, and then prepared for the Illumina HiSeq 2000 platform using a Beckmann-Coulter Roboter and the SPRIworks Fragment Library Kit I. Finally, a PCR using the KAPA PCR Amplification Kit was carried out. The libraries were then sequenced at the Columbia Genome Center to generate 30 million single-end reads of 100 bp length.

RNA-Seq analysis

Gene and isoform abundances were estimated using RSEM (with read alignment being performed by the STAR aligner) and RefSeq gene annotations (GRCh38). Differential gene expression analysis was carried out at the gene level using the edgeR and limma R packages and the Voom framework implemented therein ².

Sample Numbers

A total of 242 epithelial samples out of which 197 from primary PDAC, 26 from low-grade PanIN, and 19 from low-grade-IPMN; and a total of 159 stromal samples (matched with epithelium) out of which 124 from primary PDAC, 23 from low-grade PanIN, and 12 from low-grade IPMN were used for the study.

MUC16 expression in different subtypes of PDAC

Consensus clustering was applied on all 150 TCGA PAAD tumor samples ³ using the gene sets from Collisson et al ⁴, Moffitt et al ⁵ and Bailey et al ⁶, classifying the samples into three subtypes (classical (n=54), quasimesenchymal (n=34), or exocrine-like (n=62)), two subtypes (basal-like (n=65) or classical (n=85)) and four subtypes respectively (squamous (n=31), immunogenic (n=28), pancreatic progenitor (n=53), or aberrantly differentiated exocrine [ADEX] n=38)). Then, the Mann-Whitney rank-sum test was used for comparing MUC16 expression among subtypes.

Genomic DNA Amplification

Total DNA was extracted from T3M4 WT, WT-MUC16^{KO}, T3M4 SC, SC-MUC16^{KO} cells using Promega Wizard® genomic DNA purification kit according to the manufacturer's instructions. 100ng of total genomic DNA was amplified by PCR using Phusion™ High-Fidelity DNA Polymerase (ThermoFisher Scientific, USA) according to the manufacturer's instructions with the following primers: MUC16 FP 5'-TGGTCATTTCTGAGTGTGGAA-3', MUC16 RP 5'-CTCCACATCACCAGAGAGCA-3' (Integrated DNA Technologies, USA). PCR conditions were 39 cycles at 95°C for 5 min, 95°C for 30 sec, 60.5°C for 30 sec, 72°C for 5 min. the amplified samples were detected by 1% agarose gel electrophoresis.

SDS-Agarose electrophoresis

SDS-Agarose gel electrophoresis for MUC16 was performed as described previously ⁷ with minor modifications. Briefly, T3M4 WT, WT-MUC16^{KO}, T3M4 SC, SC- MUC16^{KO} cell lysates [RIPA lysis buffer (Thermo Scientific, Rockford, IL USA) with protease and proteinase inhibitor] were resolved on SDS (0.1%) - Agarose (2%) gel electrophoresis (50-100 µg/lane) and transferred to PVDF membrane by capillary transfer. After blocking the membranes in 5% skimmed milk in TBS-T (Tris-buffered saline and 0.1% Tween 20, pH 7.4), the membranes were probed with mouse anti-MUC16 abs, AR9.6 (Quest PharmaTech, Canada), 5B9 and 5E11 (Provided by Dr. Ulla Mandel, University of Copenhagen, Denmark), OC125 (Covance, USA), M11 (LSBio, USA) and mouse anti- α -tubulin IgG (Developmental studies hybridoma bank, Iowa, USA) overnight. The membranes were washed with TBS-T and incubated with HRP conjugated Goat-anti mouse secondary antibody (Jackson ImmunoResearch, West Grove, PA, USA). The antigen-antibody complex was detected with enhanced Chemiluminescent kit (Bio-rad, USA).

Deglycosylation of TR1.2 and MUC16

MUC16 TR1.2 glycopeptides were expressed and purified as described earlier ⁸. Deglycosylation of purified MUC16 TR1.2 glycopeptides and MUC16 positive T3M4 cell lysate was performed by using deglycosylase cocktails containing Sialidase, O-glycanase, and N-glycanase (Prozyme, Hayward, CA, USA) as per the manufacturer instructions. Deglycosylated TR1.2 glycopeptides and T3M4 cell lysates were immunoprobed with mAb AR9.6, as described above.

Isolation and purification of MUC16 from Patient's ascites fluid

MUC16 was isolated and purified from Patient's ascites fluid using standard affinity column chromatography method as described previously with modifications ⁹. Briefly, ascites fluid (10ml) was centrifuged at 1000g for 30 seconds

and mixed with protein L-resin (GenScript, USA) for 2 hours at room temperature and collected the supernatant by centrifuging at 1000g for 30 seconds. The supernatant was then passed through monoclonal antibody mAR9.6 (1mg/200 μ l) along with protein G resin (GenScript, USA) conjugated affinity chromatography column (Thermo Scientific, Rockford, IL USA). After washing out non-bound debris with PBS, the antigen was eluted from the antigen-antibody complex with acidic elution buffer (0.1M glycine-HCl, pH 2.8); neutralized with neutralization buffer (1M Tris-HCl pH 8.5); and dialyzed against PBS. The purified MUC16 antigen was used for the treatment studies.

Expression and purification of MUC16 TR1.2 in CHO cells

MUC16 TR1.2 was expressed and purified from CHO cells as described previously⁸. Briefly, CHO cells were grown in medium containing 80% EX CELL CHO Cloning Medium (Sigma) and 20% EX CELL CD CHO Fusion medium (Sigma) with 2% glutamine and 0.32 mg/mL G418 (Invitrogen). The cells were stably transfected with 3 μ g of pcDNA3- γ -MUC16 TR1.2 construct encoding for a fragment with 124 aa with a C-terminal myc tag and N- and C-terminal 6xHis tags using LipofectamineTM Transfection Reagent (ThermoFisher, USA). Positive clones were selected by screening the expression of His-tag protein with anti-His (Invitrogen, USA) or anti-MUC16 OC125 (). The secreted MUC16 TR1.2 (His-tag) was purified by cobalt column chromatography (TALONTM) as per the manufacturer's instruction.

References:

1. Radhakrishnan, P., Dabelsteen, S., Madsen, F.B., Francavilla, C., Kopp, K.L., Steentoft, C., Vakhrushev, S. Y., Olsen, J. V., Hansen, L., Bennett, E.P., et al. (2014). Immature truncated O-glycophenotype of cancer directly induces oncogenic features. *Proc Natl Acad Sci U S A.* 111, E4066-E4075.
2. Law, C.W., Chen, Y., Shi, W., Smyth, G.K. (2014). voom: Precision weights unlock linear model analysis tools for RNA-seq read counts. *Genome Biol.* 15, R29.
3. Cancer Genome Atlas Research Network. Electronic address: andrew_aguirre@dfci.harvard.edu; Cancer Genome Atlas Research Network (2017). Integrated Genomic Characterization of Pancreatic Ductal Adenocarcinoma. *Cancer Cell.* 32, 185-203.e13.
4. Collisson, E.A., Sadanandam, A., Olson, P., Gibb, W.J., Truitt, M., Gu, S., Cooc, J., Weinkle, J., Kim, G.E., akkula, L. et al. (2011). Subtypes of pancreatic ductal adenocarcinoma and their differing responses to therapy. *Nat Med.* 17, 500-503.
5. Moffitt, R.A., Marayati, R., Flate, E.L., Volmar, K.E., Loeza, G.H., Hoadley, K.A., Rashid, N.U., Williams, L.A., Eaton, S.C., Chung, A.H. et al. (2015). Virtual microdissection identifies distinct tumor- and stroma-specific subtypes of pancreatic ductal adenocarcinoma. *Nat Genet.* 47, 1168-1178.
6. Bailey, P., Chang, D.K., Nones, K., Johns, A.L., Patch, A.M., Gingras, M.C., Miller, D.K., Christ, A.N., Bruxner, T.J.C., Quinn, M.C. et al. (2016). Genomic analyses identify molecular subtypes of pancreatic cancer. *Nature.* 531, 47-52.
7. Gabriel, M., and Zentner, A. (2005). Sodium dodecyl sulfate agarose gel electrophoresis and electroelution of high molecular weight human salivary mucin. *Clin Oral Investig.* 9, 284-286.
8. Marcos-Silva, L., Narimatsu, Y., Halim, A., Campos, D., Yang, Z., Tarp, M.A., Pereira, P.J., Mandel, U., Bennett, E.P., Vakhrushev, S.Y., et al. (2014). Characterization of binding epitopes of CA125 monoclonal antibodies. *J Proteome Res.* 13, 3349-3359.
9. Lloyd, K.O., Yin, B.W., Kudryashov, V. (1997). Isolation and characterization of ovarian cancer antigen CA 125 using a new monoclonal antibody (VK-8): identification as a mucin-type molecule. *Int J Cancer.* 71, 842-850.