HAb18G/CD147 Promotes pSTAT3-Mediated Pancreatic Cancer Development via CD44s

Ling Li, Wenhua Tang, Xiaoqing Wu, David Karnak, Xiaojie Meng, Rachel Thompson, Xinbao Hao, Yongmin Li, Xiaotan T. Qiao, Jiayuh Lin, James Fuchs, Diane M. Simeone, Zhi-Nan Chen, Theodore S. Lawrence, and Liang Xu.

Laboratory of Cancer Biology, Fourth Military Medical University, Xi'an; Engineering Research Centre and Department of Cell Biology, State Key Laboratory of Cancer Biology, University of Michigan, Ann Arbor, Michigan; Cell Engineering Research Centre and Department of Cell Biology, State Key Laboratory of Cancer Biology, Fourth Military Medical University, Xi'an; Departments of Molecular Biosciences and Radiation Oncology, University of Kansas, Lawrence, Kansas; Departments of Pediatrics, College of Medicine; and Division of Medicinal Chemistry and Pharmacognosy, College of Pharmacy, Ohio State University, Columbus, Ohio

Abstract

Purpose: Signal transducer and activator of transcription 3 (STAT3) plays a critical role in initiation and progression of pancreatic cancer. However, therapeutically targeting STAT3 has failed clinically. We previously identified HAb18G/CD147 as an effective target for cancer treatment. In this study, we aimed to investigate the potential role of HAb18G/CD147 in STAT3-involved pancreatic tumorigenesis in vitro and in vivo.

Experimental Design: The expression of HAb18G/CD147, pSTAT3, and CD44s was determined in tissue microarrays. The tumorigenic function and molecular signaling mechanism of HAb18G/CD147 were assessed in in vitro cellular and clonogenic growth, reporter assay, immunoblot assay, immunofluorescence staining, immunoprecipitation, and in vivo tumor formation using loss or gain-of-function strategies.

Results: Highly expressed HAb18G/CD147 promoted cellular and clonogenic growth in vitro and tumorigenicity in vivo. Cyclophilin A (CyPA), a ligand of CD147, stimulated STAT3 phosphorylation and its downstream genes cyclin D1/survivin through HAb18G/CD147-dependent mechanisms. HAb18G/CD147 was associated and colocalized with cancer stem cell marker CD44s in lipid rafts. The inhibitors of STAT3 and survivin, as well as CD44s neutralizing antibodies suppressed the HAb18G/CD147-induced cell growth. High HAb18G/CD147 expression in pancreatic cancer was significantly correlated with the poor tumor differentiation, and the high coexpression of HAb18G/CD147-CD44s-STAT3 associated with poor survival of patients with pancreatic cancer.

Conclusions: We identified HAb18G/CD147 as a novel upstream activator of STAT3, which interacts with CD44s and plays a critical role in the development of pancreatic cancer. The data suggest that HAb18G/CD147 could be a promising therapeutic target for highly aggressive pancreatic cancer and a surrogate marker in the STAT3-targeted molecular therapies.

Introduction

Pancreatic ductal adenocarcinoma (PDCA) remains a devastating and almost uniformly lethal disease with the 5-year survival offless than 5% (1, 2). Most patients with advanced unresectable PDCA either do not respond, or respond transiently and modestly to systemic chemo/radiotherapy (3, 4). This dire clinical situation cries for our deep efforts on identifying novel therapeutic approaches.

It has been widely accepted that PDCA arises from K-ras mutations, followed by acquisition of additional epigenetic and genetic somatic alterations, including inactivation or point mutation of p16/CDKN2A, TP53, and DPC4/SMAD4 (2, 5). However, the above genetic information has not yet led to the development of effective targeted therapeutic strategies.

Signal transducer and activator of transcription 3 (STAT3) integrates signals from cytokines and growth factors into transcriptional responses in target cells. It is an important regulator of stem cell self-renewal, cancer cell survival, and...
Translational Relevance

The failure of conventional chemotherapeutic agents on survival of patients with pancreatic cancer highlights an urgent need for novel treatment strategies. STAT3 plays a critical role in pancreatic cancer initiation and progression and represents a novel therapeutic target. However, STAT3 occupies a point of convergence for many signaling pathways; blockade of existing upstream signaling to STAT3 activation is not sufficient to abrogate STAT3 activation. This study identified HAb18G/CD147 as a novel upstream activator in STAT3-mediated pancreatic tumor development by forming a signaling complex with CD44s. Furthermore, we showed that patients with pancreatic cancer with high coexpression of HAb18G/CD147-CD44s-STAT3 had poor prognosis. This information is valuable for a better understanding of the relationship between inflammation and pancreatic cancer initiation and progression induced by STAT3. Our results also suggest HAb18G/CD147 as a novel therapeutic target for highly aggressive pancreatic cancer and as a surrogate marker in clinical trials of molecular therapy targeting STAT3.

Inflammation (2, 6, 7). It was recently reported that STAT3 has critical roles in the development of PDCA, especially the initiation and progression of PDCA by controlling expression of target genes survivin, cyclin D1, and matrix metalloproteinase 7 (MMP7; refs. 5, 8, 9). In the context of acute pancreatitis and K-ras−induced pancreatic intraepithelial neoplasias (PanIN) lesions, STAT3-mediated tumor initiation are related to its ability to promote cell survival and proliferation, and to induce reprogramming of normal pancreatic epithelial cells into progenitor-like phenotype, a process assuming a proneoplastic fate (5, 8).

In addition, constitutive activation of STAT3 is frequently detected in pancreatic cancer and has been associated with a poor prognosis, and served as a therapeutic target (10). Because of lack of enzyme activity, targeting STAT3 is not easy. Inhibition of STAT3 phosphorylation/activation using monoclonal antibody or small molecules that antagonize the ligand, promotes pancreatic cancer cell growth (35, 36), and contributes to STAT3-mediated cell survival (37). These evidences promote us to investigate the role of HAb18G/CD147 in the early promotion of PDCA, especially in STAT3-involved PDCA initiation, remains largely unknown.

To explore the potential molecular targets of HAb18G/CD147, we searched the oncomine database for genes coexpressed with CD147 in pancreas (31). We observed that CD147 highly expressed in primary pancreatic cancer patients (32–34), and STAT3 is among the top listed genes that highly correlated with CD147 (Supplementary Fig. S1). It has been reported that cyclophilin A (CyPA), as a CD147 ligand, promotes pancreatic cancer cell growth (35, 36), and contributes to STAT3-mediated cell survival (37). These evidences promote us to investigate the role of HAb18G/CD147 in STAT3-mediated cell growth signaling using CyPA as a stimulus of HAb18G/CD147 activation. Typically, CD147 transmits extracellular signal by forming complexes with another membrane protein upon CyPA stimulation (38); whereas activated STAT3 promotes the transcription of target genes by phosphorylating and translation from cytoplasm into nucleus. CD44 (CD44 v3-10) has been reported to activate STAT3 signaling (39), and colocalizes with CD147 in cancer cells (40, 41), suggesting a potential role of CD44 in the CyPA-HAb18G/CD147-STAT3-mediated cell growth signaling in pancreatic tumorigenesis.

Here, we perform a systematic study to investigate the relationship among HAb18G/CD147, CD44, and STAT3 in the development of pancreatic cancer. We identified HAb18G/CD147 interacting with CD44s as a novel upstream
activator of STAT3 signaling, which plays a critical role in the development of pancreatic cancer. Our data suggest that HAb18G/CD147 could be a promising therapeutic target for highly aggressive pancreatic cancer and a surrogate marker for the STAT3-targeted molecular therapies.

Materials and Methods

Antibodies and reagents

Antibodies to phospho-STAT3 (Tyr705) and total STAT3 were purchased from Cell Signaling Technology, CD44s clone MEM-263 from Abnova, survivin from Novus, cyclin D1 and α-tubulin from Santa Cruz, goat anti-rabbit Texas Red and goat anti-mouse fluorescein isothiocyanate (FITC) from Jackson ImmunoResearch, green fluorescent protein (GFP), β-actin, goat anti-rabbit horseradish peroxidase (HRP), goat anti-mouse HRP, mouse immunoglobulin G (IgG), cholera toxin subunit B conjugates CTB-594, and geneticin (G418) from Invitrogen. Anti-mouse CD44s antibody H4C4 was obtained from the University of Iowa Developmental Studies Hybridoma Bank (Iowa, IA), and anti-mouse HAb18G/CD147 antibody HAb18 was prepared as reported (20). Recombinant human CyPA, cycloheximide, and doxycycline were purchased from Sigma. Puromycin, WP1066, and YM-155 were purchased from InvivoGen, Calbiochem, and Active Biosciences, respectively. FLLL32 were prepared as previously described (42).

Cell lines and constructs

Human pancreatic cancer cell lines and normal human embryonic lung fibroblast cell line WI-38 and embryonic kidney cell line HEK293 were obtained from the American Type Culture Collection (ATCC) and cultured in Dulbecco’s Modified Eagle Medium (DMEM; HyClone), supplemented with 10% FBS (HyClone). All cell lines from ATCC have been tested and authenticated with genotyping by ATCC and used within 6 months after receipt. The pGIPZ and pTRIPZ empty vector, CD147 pLKO.1, pGIPZ, and pTRIPZ lentiviral short hairpin RNA (shRNA) constructs were obtained from Open Biosystems. The MISSION Non-Target shRNA Control Vector (pLKO.1-NTC) was obtained from Sigma. Human CD147 cDNA was subcloned into the pEGFP-N1 expression vector (Clontech) as described (43). CD44s siRNA and control siRNA were obtained from Dharmaco. The survivin lentiviral shRNA and empty vector were obtained from OriGene. The reporter plasmid pSTAT3-luc and a pGL3-Control Vector were purchased from Promega.

Patient samples

Fresh and paraffin-blocked pancreatic tumor and adjacent nontumor tissues from patients with pancreatic cancer were obtained from University of Michigan Comprehensive Cancer Center (UMCCC; Ann Arbor, MI) Histology Core according to an Institutional Review Board-approved human protocol (H7094) and from National Engineering Center for Biochip (NECB) according to an approved human protocol. Tissue microarrays (TMA) from 193 patients were obtained from UMCCC Histology Core for analyzing HAb18G/CD147 expression in the progress of pancreatic tumorigenesis. They were constructed by two different recipient paraffin blocks: “control tissue array” and “tumor tissue array.” The “control tissue array” included 53 cases of normal pancreatic tissues and seven cases of pancreatitis; the “tumor tissue array” included two cases of PanINs, six cases of cystadenoma, 17 cases of IPMN, and 108 cases of PDCA.

For analysis of the association of HAb18G/CD147 expression with clinic-pathologic parameters, TMAs of 157 pairs of pancreatic cancer tissue and adjacent nontumor tissues (within the cancer edge of 5 cm) were obtained from NECB, the clinical, pathologic, and treatment information, together with follow-ups, and forms of consents were also obtained for these 157 patients.

Establishment of stable cell lines

The CD147 lentiviral shRNA or control shRNA was introduced into cells using FuGene6 (Roche), whereas CD147/EGFP cDNA and pEGFP control vector was transfected with HEK293 cells using Lipofectamine 2000 (Invitrogen). Knockdown or knock-in cells were selected by adding 4 to 6 μg/mL of puromycin or 1 mg/mL of G418 to the culture medium, and expression of inducible shRNA was induced in the presence of 1 μg/mL doxycycline. Silencing or increase of HAb18G/CD147 expression was verified by quantitative real-time PCR (qRT-PCR) and immunoblot analysis.

Cell growth assay

Cells were plated in 24-well plates and were counted every 24 hours for 3 to 5 days using a hemocytometer, or measuring WST-8 dye absorbance at 570 nm. A cell growth curve was drawn according to the cell numbers with the specified incubation time. For growth inhibition of STAT3/ survivin inhibitor or CD44s antibody, cells were serum-starved for 24 hours before adding 5 μmol/L FLLL32 or 0.05 to 0.1 μmol/L YM155 or 10 μg/mL H4C4 or nIG for one hour followed by 72 to 96 hours 100 nmol/L CyPA treatment. The results are shown as relative cell growth inhibition, which normalized to their individual control.

Colony formation assay

Cells were cultured in DMEM containing 10% FBS with 250 cells in each 6-well plate. Plates were maintained at 37°C in a humidified incubator for 7 to 10 days. The colonies were stained by 0.1% crystal violet and calculated under microscopy.

In vitro Matrigel invasion assay

Cells were seeded into upper chambers of 8 μm pore Transwells coated with Matrigel and then allowed to invade the Matrigel for 36 hours. Invaded cells were stained with Diff-Quik stain (Allegiance), and the invasive potential of the cells was determined by counting the number of cells that had invaded to the lower surface of the filter in 10 different areas under an inverted light microscope (Olympus BX41).
STAT3 reporter assay
Cells were seeded in 48-well plates and then transfected with 0.5 μg STAT3-luc reporter constructs or 0.5 μg of a GL3 vector control using Lipofectamine 2000. Sixteen hours after transfection, cells were treated with CyPA for 18 hours. Luciferase activity was then measured in cell lysates by a POLARstar OPTIMA Microplate Reader using the Bright-Glo Luciferase Assay System (Promega) according to the manufacturer’s protocol. For comparison, luciferase activity was normalized to β-galactosidase expression using the Beta-Glo Assay System (Promega).

Animal studies
Animal studies of tumor formation and tumor growth were performed as described previously (44). Briefly, 1 × 10⁸ cells in 0.2 mL DMEM inoculated into five- to six-week-old female athymic NCr-nu/nu nude mice subcutaneously on both flanks. Five mice with two injection sites (up and down) on each mouse were used for each type of construct (n = 10 per group). The tumor sizes and animal body weights were measured twice weekly and plotted. Tumor volume was calculated using the formula: (length × width²)/2. All animal experiments were performed according to the protocol approved by the University of Michigan Animal Care and Use Committee and in accordance with NIH guidelines.

Quantitative real-time PCR
Total RNA extraction and cDNA synthesis were carried out as previously described (45). qRT-PCR was carried out using an ABI 7700 real-time PCR system (Applied Biosystems) with gene-specific primers for HAb18G/CD147, CD44, STAT3, CyclinD1, survivin, MMP1, MMP2, MMP9, β-actin, glyceraldehyde-3-phosphate dehydrogenase (GAPDH), or 18sRNA (Supplementary Table S1). Individual genes of interest (GOI) were normalized to housekeeping genes (HKG): β-actin, 18sRNA, or GAPDH. Relative mRNA levels are presented as unit values of 2⁻ΔCt = 2⁻(Ct (HKG) – Ct (GOI)).

Immunohistochemistry
TMA staining was performed by standard immunohistochemistry procedures. To confirm the specificity of the primary antibodies, tissue sections were incubated in the absence of the primary antibodies and with control mouse IgG. The number of positively stained cells and the intensity of positive staining on epithelium and stromal cells were independently scored by two pathologists in a blinded manner. The percentage of positive stained cells was scored as: 0, 1% to 25%, 26% to 75%, and >75%. The intensity of positive immunostaining on the cells was graded by an experienced pancreatic pathologist in a blinded manner and classified into four categories: 0, 1, 2, and 3 representing no visible staining; light brown, mid-brown, and dark brown staining, respectively, with the same intensity covering more than 75% of the staining area. For statistical analysis, the stained tumor tissues were divided into two groups: the low-expression group and the high-expression group. For membranous HAb18G/CD147, only 3+ specimens were defined as high expression; for membranous CD44s and phosphorylated STAT3-positive nuclei, the samples with 1+ staining in >50% of cells or 2+ staining in >20% or all 3+ staining of cells were defined as high expression.

Immunofluorescence staining
Cells grown on chambered cover slips were either left untreated or treated with 100 mmol/L CyPA for 30 minutes. After fixedness, cells were blocked and probed with anti-HAb18G/CD147 antibody HAb18 and anti-CD44s antibody H4C4, and were then detected with fluorochrome-conjugated FITC, Texas-Red, or CTB594. Cover slips were counterstained with 4',6-diamidino-2-phenylindole (Invitrogen) for visualization of nuclei. Cell images were observed under a fluorescent microscope (Olympus).

Immunoblot and immunoprecipitation assays
Immunoblot assays were performed using standard methods (45). Cells were treated with CyPA and/or H4C4 and/or mouse IgG as required in each assay. Membranes were probed with total and phosphorylated antibodies as detailed above in Materials and Methods. Immunoprecipitation assays were performed using the Pierce Co-Immunoprecipitation (Co-IP) Kit (Thermo Fisher Scientific) according to the manufacturer’s instructions with minor modifications (45). We used 25 μg of each monoclonal antibody (complete IgG molecule) specific to either HAb18G/CD147 (HAb18) or CD44s (H4C4) for immobilization. A mouse IgG1 isotype antibody (Sigma-Aldrich) served as a negative control. Adsorbed immune complexes were washed, eluted, and then subjected to immunoblot analysis using HAb18 and H4C4.

Statistical analysis
All data shown are mean ± SEM of triplicate values from three separate experiments. *P < 0.05, **P < 0.01, and ***P < 0.001 as compared with the control group. Independent Student t tests or one-way ANOVA were used to compare the continuous variables between the two groups or more than two groups, and categorical variables were compared using the χ² test. Spearman rank correlation was conducted to analyze expression correlations among HAb18G/CD147 and CD44s. The Kaplan–Meier method and the log-rank test were used to compare overall survival, defined as the time of patients from surgery until death (patients alive were censored at the time of their last follow-up). Statistical analyses were carried out with statistical analysis software program SPSS 13.0 software (IBM) and Prism 5.0 software (GraphPad).

Results
HAb18G/CD147 is highly expressed in pancreatic cancer cell lines
To determine how HAb18G/CD147 contributes to pancreatic tumor development, we first determined the HAb18G/CD147 expression in a panel of pancreatic cancer
HAb18G/CD147 is involved in the STAT3 signaling pathway

We first determined the effects of CyPA, a natural ligand of HAb18G/CD147, on the pancreatic cancer cell growth as well as its effect on the modulation of HAb18G/CD147, STAT3, and the downstream genes cyclin D1 and survivin. We observed that CyPA induced HAb18G/CD147 protein expression in a dose- and time-dependent manner (Supplementary Fig. S5). The protein levels of pSTAT3 were significantly increased by knockdown of HAb18G/CD147 and its targets cyclin D1/survivin were also increased in a dose- and time-dependent manner upon CyPA treatment (Supplementary Fig. S5).

Next, we determined whether or not these effects are HAb18G/CD147 independent by knockdown or knock-in of HAb18G/CD147. STAT3 mRNA levels were not significantly changed (Fig. 2A) but the CyPA-induced pSTAT3 and cyclin D1/survivin protein expression were attenuated after knockdown of HAb18G/CD147 (Fig. 2B, left), supporting that the CyPA increased protein expression of pSTAT3/cyclin D1/survivin through HAb18G/CD147-dependent mechanism. Furthermore, the mRNA levels of cyclin D1 and survivin were significantly decreased by knockdown of HAb18G/CD147 in PANC-1 cells with low pSTAT3 signaling, whereas they were only slightly changed in MIA PaCa-2 cells with high pSTAT3 (Fig. 2A and Supplementary Fig. S2D), supporting that HAb18G/CD147 promoted transcription of cyclin D1/survivin in a pSTAT3-dependent manner. These results were further confirmed by knockdown of HAb18G/CD147 using pTRIPZ-inducible CD147-shRNA (Fig. 2B, right). In contrast, STAT3, cyclin D1, and survivin protein levels were significantly increased by knock-in of HAb18G/CD147 in HEK293 cells (Fig. 2B, left). Taken together, the regulation of CD147 on STAT3 appears not at the mRNA level, but at STAT3 phosphorylation. The observed colony formation inhibition by HAb18G/CD147 may be due to the decrease of pSTAT3 levels, and this effect is cell line dependent. As MIA PaCa-2 has more pSTAT3 than PANC-1, more clonogenic inhibition was observed in the MIA PaCa-2 clones with HAb18G/CD147 shRNA as compared with that in the PANC-1 clones.

Furthermore, the reporter assay showed that STAT3 transcriptional activity was significantly affected upon HAb18G/CD147 knockdown or knock-in (Fig. 2C), but no significant changes upon CyPA stimulation (data not shown). These results suggested that HAb18G/CD147 served as an upstream activator of STAT3 signaling in pancreatic cancer cells, and that HAb18G/CD147 might be more important than CyPA in STAT3 activation.

To evaluate whether STAT3 and survivin are involved in the CyPA-HAb18G/CD147-induced cell growth, we performed cell growth assay for MIA PaCa-2 cells by exposure to CyPA and either STAT3 inhibitor FLLL32/WP1066 or survivin inhibitor YM155. We observed that FLLL32/WP1066 and YM155 significantly inhibited CyPA-induced cell growth (Fig. 2D and Supplementary Fig. S5), and CyPA rescued the inhibition of cell growth by cell lines, a human pancreatic cancer primary tumor early-passage xenograft (12) and immortalized human pancreatic epithelial cell HPDE, as well as normal human embryonic lung fibroblast cell line WI-38 and embryonic kidney cell line HEK293. HAb18G/CD147 mRNA and protein expression were significantly higher in pancreatic cancer cell lines and HPDE than that in WI-38 and HEK293 cells (Supplementary Fig. S2). On the basis of the cells’ endogenous HAb18G/CD147 expression and capacity for tumor formation (44), we selected PANC-1 and MIA PaCa-2 cells for further knockdown studies and HEK293 with low endogenous HAb18G/CD147 expression for knock-in studies.

HAb18G/CD147 promotes cellular and clonogenic growth in vitro and tumor formation in vivo

To determine the potential biologic function of HAb18G/CD147 in pancreatic cancer, we adopted a loss-of-function strategy using three different lentiviral vectors: untagged plKO CD147-shRNA targeting the coding region, a GFP-tagged pGIPZ CD147-shRNA targeting the coding regions and the 3’ untranslated region (UTR), and an RFP-tagged inducible pTRIPZ CD147-shRNA targeting the coding regions and the 3’UTR. We also adopted a gain-of-function strategy to knock-in HAb18G/CD147 cDNA into HEK293 cells.

When HAb18G/CD147 expression was effectively silenced (typically 60%–80% reduction of the total; Supplementary Fig. S3 and Supplementary Table S5), which functionally validated by decrease of cell invasion (Supplementary Fig. S3), cellular and clonogenic growth was significantly decreased (Fig. 1A and B). In contrast, HAb18G/CD147 knock-in HEK293 cells with significantly increased HAb18G/CD147 expression (6.32-fold increases in mRNA levels and 3-fold increases in protein levels, comparable with the average expression in pancreatic cancer cells), have a significantly increased cell invasion (Supplementary Fig. S3) and cellular and clonogenic growth (Fig. 1A and B). These results were confirmed with pGIPZ-CD147 shRNA and pTRIPZ-inducible CD147 shRNA (Supplementary Fig. S4A and S4B).

Next, we examined the effects of HAb18G/CD147 on tumor formation in a xenograft mouse model. After knock-down of HAb18G/CD147, both tumor formation and tumor growth were significantly decreased, with a 71.8% to 87.9% reduction in tumor sizes (Fig. 1A and B). Additionally, HEK293 xenograft tumors were only slightly changed in MIA PaCa-2 cells with high pSTAT3 (Fig. 2A and Supplementary Fig. S2D), supporting that HAb18G/CD147 promotes cell invasion and tumor growth through STAT3 signaling in a pSTAT3-dependent manner. In addition, the mRNA levels of cyclin D1 and survivin were significantly decreased by knockdown of HAb18G/CD147 in PANC-1 cells with low pSTAT3 signaling, whereas they were only slightly changed in MIA PaCa-2 cells with high pSTAT3 (Fig. 2A and Supplementary Fig. S2D), supporting that HAb18G/CD147 promoted transcription of cyclin D1/survivin in a pSTAT3-dependent manner.

To determine the potential biologic function of HAb18G/CD147 on tumor formation in a xenograft mouse model. After knock-down of HAb18G/CD147, both tumor formation and tumor growth were significantly decreased, with a 71.8% to 87.9% reduction in tumor sizes (n = 10; Fig. 1C and Supplementary Fig. S4B). These results were confirmed in MIA PaCa-2 xenograft mouse model using pGIPZ CD147 shRNA (Supplementary Fig. S4D). Conversely, HAb18G/CD147 knock-in promoted tumor formation in MIA PaCa-2 cells by exposure to CyPA and either STAT3 inhibitor FLLL32/WP1066 or survivin inhibitor YM155. We observed that FLLL32/WP1066 and YM155 significantly inhibited CyPA-induced cell growth (Fig. 2D and Supplementary Fig. S5), and CyPA rescued the inhibition of cell growth by
survivin shRNA (Fig. 3A). Furthermore, the cell growth inhibition ratio of WP1066 increased up to 2-folds by knockdown of HAb18G/CD147 (Fig. 3B). These data suggest that the STAT3 and its downstream gene survivin play an important role in CyPA-HAb18G/CD147-mediated cell growth.
CD44s is involved in the activation of STAT3 by HAb18G/CD147 in lipid rafts

As shown in Fig. 3C and D, CD44s siRNA and CD44s antibody H4C4 abolished the CyPA-induced STAT3 phosphorylation and downstream signaling, indicating that CD44s is required for CyPA-HAb18G/CD147-mediated STAT3 transcription activation. CD44s protein, but not mRNA, levels were significantly affected upon CyPA stimulation (Supplementary Fig. S5D) or HAb18G/CD147 knock-down (Supplementary Fig. S6A and S6B), indicating that transcriptional regulation may not be the clue that connects HAb18G/CD147 with CD44s. Our immunoprecipitation assay showed that HAb18G/CD147 and CD44s were colocalized and significantly associated in the untreated cells (Fig. 4A and B). This association was much stronger in the HAb18G/CD147 knock-down HEK293 cells than that in HAb18G/CD147 knockdown MIA PaCa-2/A6 cells, and both were enhanced further upon CyPA stimulation. By immunofluorescence staining, we observed that both HAb18G/CD147 and CD44s were evenly located on the cell membrane before treatment, but were translocated to the GM1-enriched lipid rafts (identified by CTB-594 staining) after CyPA stimulation (Fig. 4C and Supplementary Fig. S6C). However, this translocation of CD44s in the lipid rafts was blocked after knocking down HAb18G/CD147. These results suggest that HAb18G/CD147 may colocalize and form a signaling complex with CD44s in the lipid rafts, and CyPA stimulation, though not a prerequisite, may promote the association of the two proteins.

Furthermore, we observed that the CD44s blocking antibody H4C4 attenuated the CyPA-induced cell growth.

Figure 2. CyPA-HAb18G/CD147 promotes pSTAT3-mediated cell growth. A, STAT3/cyclin D1/survivin mRNA levels in HAb18G/CD147 knockdown or knock-in cells. B, pSTAT3, tSTAT3, cyclinD1, and survivin protein levels in serum-starved HAb18G/CD147 knockdown or knock-in cells treated with or without 100 nmol/L CyPA for 30 minutes (left), and in pTRIPZ-inducible knockdown cells with or without doxycycline treatment at indicated time points (right). C, STAT3 reporter assay in HAb18G/CD147 knock-down or knock-in cells. D, cell growth assay in MIA PaCa-2 cells treated with or without STAT3 inhibitor FLLL32 (5 µmol/L), survivin inhibitor YM155 (0.05–0.1 µmol/L), and/or CyPA (100 nmol/L).
HAb18G/CD147, CD44s, and STAT3 are highly expressed in human pancreatic cancer and correlate with pancreatic cancer patients’ survival

We found that HAb18G/CD147 mRNA levels in human pancreatic tumors are significantly higher (average 4.27-fold) than that in the adjacent normal tissues (Fig. 5A). In a TMA with 193 pancreatic tissues, HAb18G/CD147 is highly expressed in only 22.6% and 14.3% of normal pancreatic tissue and chronic pancreatitis, respectively, but in 56% and 55.6% of pancreatic preneoplasia and PDCA, respectively (Fig. 5B; Table 1).

We next investigated the correlation between HAb18G/CD147 expression and clinopathological parameters in 157 cases of pancreatic cancer (Supplementary Table S2). No significant correlation exists in age, gender, tumor type, size and location, American Joint Committee on Cancer (AJCC) stage, and patient survival. But high HAb18G/CD147 expression seems to have more lymph node invasion (52.94% vs. 42.50%) and advanced tumor–node–metastasis (TNM) stage (51.35% vs. 37.5%). Moreover, high HAb18G/CD147 expression was found to be significantly associated with a poor tumor differentiation (51.43% vs. 15.65%, P < 0.0001). Taken together, being upregulated in both pancreatic preneoplasia and PDCA, and being correlated to poor tumor differentiation and advanced TNM stage, HAb18G/CD147 may have a role in pancreatic tumor development.

To investigate whether HAb18G/CD147-mediated tumor growth and CD44s-pSTAT3 expression are associated in patient samples, we analyzed the coexpression of CD44s-pSTAT3 with HAb18G/CD147 in 157 pancreatic cancer tissues. As indicated, high HAb18G/CD147 expression was significantly correlated with high CD44s expression (Spearman r = 0.4961, P < 0.001; Supplementary Fig. S6D and S6E), but not with high pSTAT3 nuclear positivity (Table 2).

With regard to the clinical pathologic factors, patients with high expression of HAb18G/CD147 and CD44s had poor tumor differentiation (P = 0.0001; Supplementary Table S3) and a lower median survival, as compared with patients with low expression of HAb18G/CD147 and CD44s, although the latter lacks statistical difference (P = 0.311; Fig. 5C, top).

Moreover, patients with high expression of all three genes had a higher incidence of poor tumor differentiation (60.87% vs. 7.69%, P = 0.002) and mortality (60% vs. 28.57%, P = 0.218), as well as a 75% survival of 4 months, as compared with that of 10 months for patients with low expression of the three genes (HR of death = 3.024; 95% confidence interval, 0.642–15.983, P = 0.12; Fig. 5C and Supplementary Table S4). These results suggest that HAb18G/CD147-CD44s-pSTAT3 association might be used as a prognosis marker for pancreatic cancer.
Discussion

In this study, we found that HAb18G/CD147 was highly expressed in pancreatic cancer cell lines, chronic pancreatitis, preneoplasia, and PDCA. Knockdown of HAb18G/CD147 significantly inhibited the tumor cell invasion, cellular and clonogenic growth in vitro and reduced tumor formation and tumor growth in xenograft mouse model. Moreover, we showed that HAb18G/CD147 and CD44s are involved in the activation of STAT3 signaling pathway. Finally, we demonstrated that patients with high expression of HAb18G/CD147, CD44s, and pSTAT3 had higher mortality and poorer tumor differentiation, and poorer survival, as compared with the patients with low expression of these three genes. These data support that HAb18G/CD147 plays a novel oncogenic role in pancreatic tumor development, besides its well-known role in regulating MMP-related metastasis.

We previously showed that HAb18G/CD147 plays an important role in liver tumor metastasis under different experimental and clinical conditions (13, 20). In this study, we showed that HAb18G/CD147 plays a role in early promotion of pancreatic cancer. HAb18G/CD147 antibody has been used in clinical practice for liver cancer therapy, and other CD147 antibodies are in preclinical development for pancreatic cancer (29, 30, 46), suggesting the potential clinical application of HAb18G/CD147 antibody for patient with PDCA.

Both HAb18G/CD147 activator CyPA and STAT3 were related to inflammation and cancer, suggesting a potential link between inflammation and pancreatic tumor initiation via CyPA-HAb18G/CD147-STAT3 signaling. STAT3, as a mediator of inflammation-associated processes, plays a critical role in PDCA initiation and progression and has been under active investigation as a potential target for PDCA therapy (5, 8, 9). However, STAT3-targeted anticancer drug, either anti-EGFR antibodies or small-molecule tyrosine kinase inhibitors, only showed a limited efficacy (10). As STAT3 occupies a point of convergence for many signaling pathways, blockade of existed upstream signaling to STAT3 activation may not sufficiently abrogate STAT3. In
this study, we showed that CyPA induced STAT3 phosphorylation through HAb18G/CD147-dependent mechanism, suggesting that HAb18G/CD147 is a real upstream regulator of STAT3 activation. Pancreatic cancer cells with high HAb18G/CD147 levels and/or activity had higher active STAT3 signaling, on which the cells depend for survival and

<table>
<thead>
<tr>
<th>Table 1. HAb18G/CD147 expression in pancreatic tissues</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
</tr>
<tr>
<td>Normal</td>
</tr>
<tr>
<td>Pancreatitis</td>
</tr>
<tr>
<td>Preneoplasms</td>
</tr>
<tr>
<td>PanIN</td>
</tr>
<tr>
<td>CYSTADEMA</td>
</tr>
<tr>
<td>IPMN</td>
</tr>
<tr>
<td>PDCA</td>
</tr>
</tbody>
</table>

Boldface: compared with normal, the significance of PanIN was 0.963 and that of CYSTADEMA was 0.335.

A HAb18G/CD147 expression levels are scored as: 0 (no staining), 1 (light staining), 2 (intermediate staining), and 3 (intense staining).

*Estimated by χ² test as compared with normal tissues by dividing into low- and high-expression group. Low-expression group are cases with staining intensity scores of 0 to 2; high-expression group are cases with staining intensity scores of 3.
proliferation, much like the so-called “oncogene addiction.” Actually, CyPA has previously been reported to regulate STAT3 tyrosine phosphorylation and nuclear translocation (37). We also showed that HAb18G/CD147 promotes tumor growth by regulating the expression of survivin and cyclin D1. This result is consistent with previous reports that STAT3 promotes tumor early promotion by controlling the transcription of antiapoptotic gene survivin and proliferative gene cyclin D1 (8). These data suggest that HAb18G/CD147 may exert its tumor-promoting function by activating the STAT3 phosphorylation, indicating a novel role for HAb18G/CD147 in the STAT3-mediated tumor early promotion in PDCA. It has been reported that the expression of EMMPRIN (CD147) may be modulated by STAT3 ODN (47), indicating a positive feedback loop between HAb18G/CD147 and STAT3. In other word, CyPA-HAb18G/CD147 activates STAT3 phosphorylation, and then pSTAT3 promotes HAb18G/CD147 transcription. Thus, the CyPA-HAb18G/CD147-STAT3-mediated growth-promoting signaling could be amplified, in which case STAT3 inhibitors may have better effects for cancer cells with high expression of HAb18G/CD147. In a rationally designed clinical trial of STAT3-targeted cancer therapy, the HAb18G/CD147 level in patients with cancer should also be considered. This will help us to select a patient subgroup that is more likely to respond to the molecularly targeted therapy. HAb18G/CD147 may thus become an important surrogate marker in clinical trials of molecular therapies targeting STAT3.

HAb18G/CD147 expression is highly correlated with CD44s expression in our analysis. Patients with HAb18G/CD147high/CD44s high pSTAT3 high cytoplasmic localization show high mortality, implying a potential link of HAb18G/CD147 to tumor initiation by associating with the cancer stem cell (CSC) marker CD44s (48, 49), and by regulating the CSC signaling molecule STAT3 (50). CD147 has been shown to be a CSC marker (48, 49), and by regulating the CSC signaling molecule STAT3 (50). CD147 has been shown to be a CSC marker (48, 49), and by regulating the CSC signaling molecule STAT3 (50). However, HAb18G/CD147-CD44s-STAT3 indicates poor prognosis with the CSC marker CD44s, and that coexpression of HAb18G/CD147 with CD44s, but not CD44v, plays a role in pancreatic tumor development. Furthermore, HAb18G/CD147 and CD44s distribute together in the specific signaling platform, lipid rafts, which facilitate efficient downstream signal transduction. The distribution and activity of CD44s depend on CyPA-HAb18G/CD147, which is consistent with a previous report indicating that HAb18G/CD147 membrane localization is not affected by hyaluronic oligomers that antagonize the hyaluronan–CD44 interaction (51). Therefore, we propose that upon CyPA stimulation, HAb18G/CD147 first clusters and recruits CD44s in the lipid rafts to form a signaling complex, and then promotes STAT3 phosphorylation and cyclin D1/survivin transcription, finally leads to cell survival and cell-cycle progression (Fig. 5D).

In conclusion, our study suggests that HAb18G/CD147 is a novel upstream activator in STAT3-mediated pancreatic tumor development by forming signaling complex with the CSC marker CD44s, and that coexpression of HAb18G/CD147-CD44s-STAT3 indicates poor prognosis in patients with pancreatic cancer. This information will be valuable for a better understanding of the relationship between inflammation and pancreatic cancer initiation and progression induced by STAT3; in addition, our results establish HAb18G/CD147 as a novel therapeutic target for highly aggressive pancreatic cancer and as a surrogate marker in clinical trials of molecular therapies targeting STAT3.

**Disclosure of Potential Conflicts of Interest**
No potential conflicts of interest were disclosed.

**Authors’ Contributions**

Conception and design: L. Li, Z.-N. Chen, T.S. Lawrence, L. Xu
Acquisition of data (provided animals, acquired and managed patients, provided facilities, etc.): L. Li, W. Tang, D. Karnak, X. Wu, X. Meng, R. Thompson, X. Hao, X.T. Qiao, J. Fuchs, L. Xu
Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis): L. Li, W. Tang, X. Meng, X.T. Qiao, D.M. Simeone, T.S. Lawrence, L. Xu

Table 2. Spearman correlation coefficients for the correlation among expression of HAb18G/CD147, STAT3, and CD44s

<table>
<thead>
<tr>
<th>Gene expression</th>
<th>HAb18G/CD147</th>
<th>Nuclear pSTAT3</th>
<th>Cytoplasm pSTAT3</th>
<th>CD44s</th>
</tr>
</thead>
<tbody>
<tr>
<td>HAb18G/CD147</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td></td>
</tr>
<tr>
<td>Nuclear pSTAT3</td>
<td>−0.035</td>
<td>−0.098</td>
<td>0.415a</td>
<td>0.496a</td>
</tr>
<tr>
<td>Cytoplasm pSTAT3</td>
<td>−0.098</td>
<td>0.415a</td>
<td>—</td>
<td>0.019</td>
</tr>
<tr>
<td>CD44s</td>
<td>0.496a</td>
<td>0.165b</td>
<td>0.019</td>
<td>—</td>
</tr>
</tbody>
</table>

*P < 0.001.
*P < 0.05.
Writing, review, and/or revision of the manuscript: L. Li, R. Thompson, X.T. Qiao, D.M. Simonce, Z.-N. Chen, T.S. Lawrence, L. Xu

Administrative, technical, or material support (i.e., reporting or organizing data, constructing databases): L. Li, X. Meng, X. Hao, Y. Li, X.T. Qiao, J. Lin, L. Xu

Study supervision: L. Li, Z.-N. Chen, L. Xu

Acknowledgments

The authors thank Dr. Susan Harris for help with the editing of the manuscript, Drs. Thomas Godano and Dafydd Thomas and Michelle Vinco in the UMCCC Tissue Core for providing tissues and TMAs and help on immunohistochemistry staining. The authors also thank the UMCCC Unit of Laboratory Animal Medicine (ULAM) for help with the animal experiments.

References

6. Li N, Grivennikov SI, Karin M. The unholy trinity: in
7. Yu H, Pardoll D, Jove R. STATs in cancer in
8. Perez-Mancera PA, Guerra C, Barbacid M, Tuveson DA. What we have

Grant Support

This work was supported in part by U.S. NIH grants R01 CA121830 S1 (to L. Xu), R01 CA134655 (to L. Xu), and 5P30 CA46592 (University of Michigan Cancer Center Support Grant), Kansas Bioscience Authority Rising Star Award (to L. Xu), China National Science and Technology Major Project 2013ZX09301030 (to Z.-N. Chen), and National Basic Research Program 2009CB521704 (to Z.-N. Chen).

The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked advertisement in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

Received March 11, 2013; revised September 11, 2013; accepted October 3, 2013; published OnlineFirst October 16, 2013.


HAb18G/CD147 Promotes pSTAT3-Mediated Pancreatic Cancer Development via CD44s

Ling Li, Wenhua Tang, Xiaoqing Wu, et al.

Cite article

Updated version
Access the most recent version of this article at: doi:10.1158/1078-0432.CCR-13-0621

Supplementary Material
Access the most recent supplemental material at: http://clincancerres.aacrjournals.org/content/suppl/2013/10/16/1078-0432.CCR-13-0621.DC1

Cited articles
This article cites 50 articles, 14 of which you can access for free at: http://clincancerres.aacrjournals.org/content/19/24/6703.full.html#ref-list-1

E-mail alerts
Sign up to receive free email-alerts related to this article or journal.

Reprints and Subscriptions
To order reprints of this article or to subscribe to the journal, contact the AACR Publications Department at pubs@aacr.org.

Permissions
To request permission to re-use all or part of this article, contact the AACR Publications Department at permissions@aacr.org.