Inhibition of Class I Histone Deacetylases Abrogates Tumor Growth Factor β Expression and Development of Fibrosis during Chronic Pancreatitis

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ABSTRACT

Pancreatic fibrosis is the hallmark of chronic pancreatitis, a highly debilitating disease for which there is currently no cure. The key event at the basis of pancreatic fibrosis is the deposition of extracellular matrix proteins by activated pancreatic stellate cells (PSCs). Transforming growth factor β (TGFβ) is a potent profibrotic factor in the pancreas as it promotes the activation of PSC; thus, pharmacologic interventions that effectively reduce TGFβ expression harbor considerable therapeutic potential in the treatment of chronic pancreatitis. In this study, we investigated whether TGFβ expression is reduced by pharmacologic inhibition of the epigenetic modifiers histone deacetylases (HDACs). To address this aim, chronic pancreatitis was induced in C57BL/6 mice with serial injections of cerulein, and the selective class 1 HDAC inhibitor MS-275 was administered in vivo in a preventive and therapeutic manner. Both MS-275 regimens potently reduced deposition of extracellular matrix and development of fibrosis in the pancreas after 4 weeks of chronic pancreatitis. Reduced pancreatic fibrosis was concomitant with lower expression of pancreatic TGFβ and consequent reduced PSC activation. In search of the cell types targeted by the inhibitor, we found that MS-275 treatment abrogated the expression of TGFβ in acinar cells stimulated by cerulein treatment. Our study demonstrates that MS-275 is an effective antifibrotic agent in the context of experimental chronic pancreatitis and thus may constitute a valid therapeutic intervention for this severe disease.

Introduction

Chronic pancreatitis is defined as a progressive inflammation of the pancreas, resulting in development of organ fibrosis, which is at the core of the disease pathophysiology. This progressive condition is characterized by irregular sclerosis with focal, segmental, or diffuse destruction of the parenchyma. Consequently, gradual loss of exocrine and endocrine cellular components leads to pancreatic insufficiency and eventually diabetes, which are associated with considerable morbidity, reduction of quality of life, and reduction of life expectancy (reviewed in DiMagno and DiMagno, 2013, 2016).

Fibrosis is characterized by excessive production and deposition of extracellular matrix (ECM) components in the pancreatic parenchyma, produced mainly by resident pancreatic stellate cells (PSCs). In response to organ injury, profibrogenic factors are released and activate PSCs, a process characterized by phenotypical cell alteration, proliferation, and ECM protein synthesis.

Despite advances in chronic pancreatitis research, to date, the complex cellular and signaling mechanisms that drive the fibrotic process are not yet completely elucidated. This limited knowledge explains why therapeutic approaches to counteract the development of organ fibrosis are not currently available and management of chronic pancreatitis remains a clinical challenge.

In this study, we evaluated whether administration of MS-275 (also known as entinostat), a selective inhibitor of class 1 histone deacetylases (HDACs), counteracts the development of pancreatic fibrosis using the widespread murine model of cerulein-induced chronic pancreatitis. The rationale for this approach was 3-fold: 1) development of fibrosis activates a substantial gene regulation, which is prominently orchestrated by epigenetic mechanisms (McDonnell et al., 2014; Weigel et al., 2015; Yang and Schwartz, 2015; Moran-Salvador and Mann, 2017); 2) HDACs are critical epigenetic regulators, and expression of class 1 HDACs is significantly upregulated during the course of chronic pancreatitis (Bombardo et al., 2017); and 3) pharmacologic inhibitors of

ABBREVIATIONS: α-SMA, α-smooth muscle actin; ECM, extracellular matrix; HDAC, histone deacetylase; PCR, polymerase chain reaction; PSC, pancreatic stellate cell; TGFβ, transforming growth factor β.
HDAC activity, originally developed as anticancer agents, are currently being investigated for their antifibrotic properties in different fibrotic diseases (recently reviewed in Pang and Zhuang, 2010; Royce et al., 2014; Chen et al., 2015; Schuetze et al., 2016).

Materials and Methods

Animal Experiments. All animal treatments were performed in accordance with Swiss federal animal regulations and approved by the cantonal veterinary office of Zurich. All studies involving animals were carried out in accordance with the Guide for the Care and Use of Laboratory Animals as adopted and promulgated by the U.S. National Institutes of Health. Mice used in this study were adult 6- to 10-week-old wild-type C57BL/6 mice in a weight range of 25–30 g (Envigo Laboratories, Horst, The Netherlands). Animals were kept under standardized conditions under 12-hour light/dark cycles, with food and water available ad libitum. Groups of 4 to 5 mice were kept in standard individually ventilated cages in a specific pathogen-free facility. Food and water were provided ad libitum. Only male mice were used in this study.

Chronic pancreatitis was induced via six intraperitoneal injections of cerulein (50 µg/kg) administered hourly every 2nd day for up to 6 weeks. Control animals received 0.9% NaCl injections. MS-275 (Selleckchem, Houston, TX) was injected intraperitoneally at 20 mg/kg every 2nd day for 2 weeks, starting concomitantly (preventive regimen) or 1 week after the beginning of cerulein injections (therapeutic regimen). The concentration of MS-275 was chosen based on previously published in vivo studies using the inhibitor in mice (Dalgaard et al., 2008; Nguyen et al., 2008; Murphy et al., 2014; Bombardo et al., 2017). Control animals received 10% dimethylsulfoxide injections. Intraperitoneal injections were alternated between the left and right sides of the abdomen. Mice were examined throughout the development of pancreatitis, and their health status was recorded every 2nd day on a score sheet. Pancreatitis models used in this study generated only a mild form of the disease. Animal weight loss did not exceed 10%, and no mortality was observed. After deep terminal anesthesia with isoflurane, mice were euthanized via cardiac puncture exsanguination. Groups of five animals were tested for each experiment. Animals were assigned randomly to different experimental groups for all in vivo studies. Data collection and evaluation of all in vivo and in vitro experiments were performed blinded to group identity.

Mammalian Cell Cultures. Cell culture reagents were from Gibco-BRL. Rat AR42J cells were maintained in Kaighn’s F-12 medium with 20% fetal bovine serum, supplemented with 50 U/ml penicillin and 50 µg/ml streptomycin, and maintained at stable condition of 37°C in a 5% CO2 atmosphere. Cells were seeded in six-well plates, stimulated with 10 nM cerulein for 4 hours, and lysed in the plates for RNA extraction and real-time polymerase chain reaction (PCR) analysis.

Primary acini were isolated according to Algul et al. (2007) from 6-week-old Wistar male rats from Charles River, Germany. Acini were preincubated with 1 µM MS-275 for 30 minutes and stimulated with 0.1 nM cerulein for 30 minutes in the presence of 1 µM MS-275. At the end of the treatment, cells were lysed in the plates for RNA extraction and real-time PCR analysis.

Immunohistochemistry. Pancreas specimens were embedded in paraffin for histologic analyses, as previously described (Silva et al., 2011). H&E and Masson’s trichrome staining were performed according to routine procedures. Microscopy analyses were performed on a wide-field Nikon Eclipse Ti (Amsterdam, The Netherlands). Quantification of labeled cells was performed in at least 10 randomly selected high-power fields (×200) per slide using the NIS Elements BR Analysis (Nikon, Amsterdam, The Netherlands) and Cell*P analysis software (Olympus, Tokyo, Japan).

Western Blotting. Twenty milligrams of pancreatic tissue was homogenized in radioimmunoprecipitation (RIPA) assay buffer containing a protease inhibitor cocktail (Roche Diagnostics, Mannheim, Germany). Protein concentrations were determined by a Bradford protein assay (BioRad, Hercules, CA). Twenty micrograms of proteins was resolved by SDS-PAGE electrophoresis and blotted onto nitrocellulose membranes using a V3 Western Workflow system (BioRad) according to the manufacturer’s protocols.

Membranes were incubated with primary antibodies overnight at 4°C. Primary antibodies used in this study were: mouse anti-α-smooth muscle actin (α-SMA, Dako, Glostrup, Denmark); rabbit anti α-tubulin (ab52894; Abcam, Cambridge, UK); rabbit anti-phospho-Smad3 (Ser423/425) (Cell Signaling, Danvers, MA); and rabbit anti-GAPDH (Santa Cruz Biotechnology, Dallas, TX).

Nuclear Protein Extraction and HDAC Activity. Nuclear proteins were extracted from 20 mg of pancreatic tissue with the EpiQuik Nuclear Extraction Kit (Epigentek Group Inc, Mountain View, CA), and HDAC activity was measured in the nuclear extracts with the fluorimetric EpiQuik HDAC activity/inhibition assay kit (Epigentek Group Inc.), according to the manufacturer’s instructions.

Transcript Analyses. Total RNA was extracted from pancreatic tissue and acinar explants as described previously (Graf et al., 2002) and reverse-transcribed with qScript cDNA SuperMix (Quanta Biosciences, Beverly, MA). Gene expression was measured by real-time PCR on a 7500 Fast Real-Time PCR System (Applied Biosystems, Carlsbad, CA) using Taqman probes (Applied Biosystems). Transcript levels were normalized using 18S RNA as a reference and expressed as ΔΔCt relative to the value of control animals or as ΔCt.

Statistical Analyses. Every group of mice compared in the different experimental conditions was comprised 5 animals. Data are expressed as means ± standard deviation. Population characteristics were compared among treatment groups using an unpaired, two-tailed Student’s t test when comparing two experimental conditions or one-way analysis of variance, followed by Dunnett’s post-hoc test when comparing more than two experimental conditions. Holm-Bonferroni correction for multiple comparisons was used to keep the family-wise error rate of dependent variables at 5%. Analyses were performed using GraphPad Prism 4.0c (GraphPad Software, Inc., San Diego, CA).

Results

Development of Fibrotic Response during Chronic Pancreatitis Correlates with Increased Levels of HDAC Expression. To investigate the role of HDAC in the development of pancreatic fibrosis after induction of chronic pancreatitis, we first performed a time-course analysis after induction of the disease to determine the kinetics of the fibrotic response. Histologic evaluation of mice harvested after 2, 4, and 6 weeks of cerulein treatment revealed progressive damage of pancreatic parenchyma and cell infiltration (Fig. 1A) and pronounced ECM deposition (Fig. 1B). Quantification of fibrotic parameters showed increased expression of collagen isoforms (Fig. 1C) and collagen deposition in the pancreas (Fig. 1D). Development of pancreatic fibrosis is mediated by activated pancreatic stellate cells (PSCs), which constitute the predominant source of ECM proteins, including collagens and fibroactin. Activation of PSCs, detected by α-smooth muscle actin (α-SMA) expression, reached a maximum after 4 weeks of pancreatitis (Fig. 1E), thus mirroring the kinetics of collagen expression. Expression of profibrotic TGFβ isoforms, the main activators of PSC, and TGFβ receptor II, critical for the development of pancreatic fibrosis (Yoo et al., 2005), also increased in a similar pattern during
the development of organ fibrosis (Fig. 1F). In addition, pancreatic expression of inflammatory components followed similar kinetics (Supplemental Fig. 1) (Bombardo et al., 2017). Based on these results showing maximal levels of fibrosis after 4 weeks of pancreatitis, we focused on this time point for further analyses.

**Inhibition of Class 1 HDAC with MS-275 Reduces the Development of Fibrosis after Induction of Chronic Pancreatitis.** We recently showed that gene expression levels of class 1 HDACs were upregulated during chronic pancreatitis (Bombardo et al., 2017). This finding was confirmed by increased HDAC enzymatic activity (Fig. 2A) in
Fig. 2. Preventive and therapeutic administration of MS-275 reduces the development of fibrosis during chronic pancreatitis. (A) Total HDAC activity detected in pancreatic nuclear extract in control and mice treated with cerulein (Cer) for 4 weeks mice. (B) Schematic representation of 2 weeks of preventive (MS + Cer) and therapeutic (Cer + MS) MS-275 regimens during induction of chronic pancreatitis. Cer was administered on alternate days over 4 weeks. MS-275 was administered on alternate days over 2 weeks. (C) H&E staining of pancreata after 4 weeks of chronic pancreatitis after preventive and therapeutic MS-275 regimens. (D) Masson’s trichrome staining of pancreata showing reduced collagen deposition (green) after 4 weeks of
pancreatic nuclear proteins after 4 weeks of cerulein treatment. To test whether class 1 HDAC upregulation was functionally linked to the development of fibrosis, we treated mice with the selective class 1 inhibitor MS-275, which we previously showed to potently inhibit HDAC activity in the pancreas (Bombardo et al., 2017). The inhibitor was administered for 2 weeks in a preventive manner, starting concomitantly with the first cerulein injection. Alternatively, we administered the inhibitor in a therapeutic manner, starting 1 week after the induction of pancreatitis. Animals were harvested after 4 weeks of chronic pancreatitis according to the regimens depicted in Fig. 2B. Both modalities of MS-275 treatment resulted in a trend of reduced expression of selected HDAC isoforms (Supplemental Fig. 2) and better preservation of pancreatic parenchyma (Fig. 2, C and D). This result was further confirmed by reduced expression of collagen isoforms (Fig. 2E) and lower collagen deposition (Fig. 2F) after MS-275 administration. As collagen is deposited mainly by activated PSCs, we next evaluated whether PSC activation was limited in the presence of MS-275. Expression of α-SMA, a key hallmark of PSC activation, was lower upon MS-275 treatment at both RNA (Fig. 3A) and protein levels (Fig. 3B). These data revealed that both preventive and therapeutic regimens of MS-275 were effective in reducing PSC activation and consequently limiting the development of fibrotic processes on induction of chronic pancreatitis.

Inhibition of Class 1 HDAC with MS-275 Reduces TGFβ Expression after Induction of Chronic Pancreatitis. Activation of PSCs is initiated by damaged acinar cells and reinforced by already activated PSCs in a paracrine and autocrine manner through the synthesis and secretion of profibrotic mediators (Apte et al., 2011). Thus, we investigated whether the reduced fibrosis observed upon MS-275 treatment was linked to reduced expression of these factors. In support of this hypothesis, gene expression levels of profibrotic mediators (Apte et al., 2011). Thus, we investigated whether the reduced fibrosis observed upon MS-275 treatment was linked to reduced expression of these factors. In support of this hypothesis, gene expression levels of profibrotic TGFβ1–3 were lower in the presence of the inhibitor (Fig. 4A). Consequent to the decreased production of TGFβ isoforms, activation of TGFβ signaling was attenuated, as shown by a trend of reduced C-terminal phosphorylation that is required for activation of Smad3, an initiating event of the intracellular cascade resulting upon engagement and dimerization of TGFβ receptor complex (Fig. 4B). Interestingly, TGFβ receptor 2 was also upregulated following induction of pancreatitis; however, its expression levels were not reduced after MS-275 treatment (Fig. 4C).

Inhibition of Class 1 HDAC with MS-275 Reduces TGFβ Expression in Pancreatic Acinar Cells. Previous works have shown that TGFβ is synthesized in activated PSC, thus acting through autocrine loops (reviewed in Apte et al., 2011); however, in our study, we investigated whether the synthesis of TGFβ upstream of PSC activation is inhibited by MS-275. Specifically, we asked whether: 1) acinar cells upregulate TGFβ expression after cerulein-induced pancreatitis and 2) MS-275 selectively interferes with this gene expression. In support of this hypothesis, we detected early upregulation of TGFβ1 in the pancreas 24 hours after cerulein treatment (Fig. 5A). Furthermore, in a new set of experiments, upregulation of TGFβ1-2 isoforms and TGFβ receptor 2 was observed in primary acini isolated 24 hours after cerulein treatment in vivo (Fig. 5B), suggesting that acini respond to the initial injury by upregulating TGFβ signaling components. We then tested whether TGFβ upregulation in acinar cells was a direct effect of cerulein administration or rather stimulated by stromal cells. AR42J acinar cells treated in vitro with cerulein upregulated TGFβ1 levels (Fig. 5C). Similarly, upregulation of TGFβ1 was observed when primary pancreatic acini were isolated and stimulated in vitro with cerulein (Fig. 5D). These data suggest that cerulein induced TGFβ1 gene expression in a cell-autonomous manner independent from the presence of stromal cells. Importantly, pretreatment

chronic pancreatitis after preventive and therapeutic MS-275 regimens. (E) qPCR of α-SMA expression in pancreata after 4 weeks of chronic pancreatitis after preventive and therapeutic MS-275 regimens. (B) Western blot quantification of α-SMA in pancreata after 4 weeks of chronic pancreatitis after preventive and therapeutic MS-275 regimens. Band intensity values were normalized using tubulin as a loading control. Results are average ± S.D. (n = 5), *P < 0.05.

Fig. 3. Preventive and therapeutic administration of MS-275 reduces the activation of pancreatic stellate cells during chronic pancreatitis. (A) qPCR of α-SMA expression in pancreata after 4 weeks of chronic pancreatitis after preventive and therapeutic MS-275 regimens. (B) Western blot quantification of α-SMA in pancreata after 4 weeks of chronic pancreatitis after preventive and therapeutic MS-275 regimens. Band intensity values were normalized using tubulin as a loading control. Results are average ± S.D. (n = 5), *P < 0.05.

Fig. 4. Preventive and therapeutic administration of MS-275 reduces the activation of pancreatic stellate cells during chronic pancreatitis. (A) qPCR of α-SMA expression in pancreata after 4 weeks of chronic pancreatitis after preventive and therapeutic MS-275 regimens. (B) Western blot quantification of α-SMA in pancreata after 4 weeks of chronic pancreatitis after preventive and therapeutic MS-275 regimens. Band intensity values were normalized using tubulin as a loading control. Results are average ± S.D. (n = 5), *P < 0.05.
with the HDAC inhibitor MS-275 abrogated TGFβ1 induction in both AR42J cells and primary acini (Fig. 5, C and D).

Furthermore, cerulein treatment increased expression of and HDAC1, but not HDAC2 and 3, in isolated acini (Fig. 5E), further suggesting that TGFβ1 expression in acinar cells is promoted by HDAC activity. Expression of TGFβ2 was much lower than TGFβ1 and not regulated in this experimental setting in both cell types (Supplemental Fig. 3, A and B).

Discussion

TGFβ is a potent fibrogenic factor that plays a pivotal role in the development of fibrosis during chronic pancreatitis (Menke et al., 1997; Yoo et al., 2005; He et al., 2009; Li et al., 2016). One of the main effects exerted by TGFβ is activation of PSCs from a quiescent state to a myofibroblast-like phenotype (reviewed in Apte et al., 2011). In recent years, activated PSCs have attracted increasing attention as major mediators of pancreatic fibrosis during chronic pancreatitis as they not only mediate the development of fibrosis by producing ECM proteins, but they also amplify the fibrotic response in an autocrine and paracrine manner by secreting fibrogenic factors, including TGFβ (Kruse et al., 2000). In the present study, we discovered that activity of HDACs in the pancreas is functionally linked to the development of fibrosis during chronic pancreatitis, thus providing a potential therapeutic target to counteract this disease. This hypothesis was further tested in in vivo experiments where the selective inhibitor of class 1 HDACs MS-275 was administered in a preventive or therapeutic manner during chronic pancreatitis. In both regimen types, we observed a striking inhibition of pancreatic fibrosis and increased preservation of pancreatic parenchyma, suggesting that MS-275 exerts an antifibrotic effect, even when administered after commencement of the disease.

At the cellular level, reduced fibrosis detected upon MS-275 treatment was likely the result of reduced TGFβ expression, leading to reduced activation of PSCs. An important question arising from these data is the identity of the cells whose TGFβ production is targeted by the inhibitor. Using in vitro experiments with isolated acinar cells, we found that short-term incubation with cerulein was sufficient to stimulate TGFβ expression in these cells. The fact that MS-275 treatment potently reduced cerulein-stimulated TGFβ expression suggests that acinar cells are indeed a direct target of the inhibitor and contribute to the phenotype observed in vivo. In this regard, it would be important to explore further the temporal regulation of TGFβ isoform expression in acinar cells to dissect the dynamic of their contribution to the development of pancreatic fibrosis.

It is known that acinar cells are not the only source of TGFβ in the pancreas, as previous studies reported the presence of TGFβ1 mRNA in stromal cells, including PSC, upon induction of pancreatitis (Muller-Pillasaki et al., 1999). In this regard, it is important to mention that TGFβ synthesis in nonacinar
cells may also depend on HDAC activity. In fact, treatment of isolated PSCs with the pan-HDAC inhibitor sodium valproate inhibits TGFβ expression and collagen synthesis in these cells (Bülow et al., 2007). Furthermore, another possible source of TGFβ production is inflammatory cells, which are recruited to the pancreas during the development of pancreatitis. This is particularly interesting as we recently demonstrated that MS-275 treatment effectively reduced the levels of inflammation during the course of acute and chronic pancreatitis (Bombardo et al., 2017). Crosstalk between PSCs and distinct leukocyte populations, including macrophages, promotes PSC activation and fibrosis during chronic pancreatitis (Xue et al., 2015). Future studies using coculture of acinar cells, PSC, and leukocytes are warranted to dissect the contribution of the individual cell types in the production of TGFβ upon treatment with MS-275 and the effect on PSC activation.

Although it is possible that reduced inflammation upon MS-275 administration leads to reduced fibrosis during chronic pancreatitis, a recent study revealed that development of inflammation and fibrosis is two independent and, accordingly, not causal events in this disease. Specifically, using transgenic mice deficient in Cxcr2, the authors observed almost complete ablation of inflammatory cell infiltration upon chronic pancreatitis; however, this limited inflammatory reaction did not prevent PSC activation; consequently, fibrosis levels were comparable in transgenic and wild-type control mice (Steele et al., 2015). This striking example implies that signaling molecules derived from inflammatory cells may play a minor role in the development of pancreatic fibrosis during chronic pancreatitis.

Collectively, our results integrate with the current body of evidence demonstrating the crucial role of HDACs in the development of fibrotic diseases. In this regard, compelling
evidence demonstrates that HDAC activity is necessary for activation of hepatic stellate cells in vitro (reviewed in Chen et al., 2015). The requirement of HDAC activity in driving myofibroblastic differentiation and ECM protein synthesis was also observed in different fibroblast populations present in skin, lung, and kidney (Glenisson et al., 2007; Yoshikawa et al., 2007; Guo et al., 2009). Moreover, the use of different HDAC inhibitors showed beneficial effects in the treatment of hepatic, renal, cardiac, and pulmonary fibrosis in vivo (Kee et al., 2013; Liu et al., 2013; Van Beneden et al., 2013; Khan and Jena, 2014; Nural-Guven et al., 2014; Chen et al., 2015; Choi et al., 2015; Korfei et al., 2015). This finding suggests that epigenetic mechanisms controlled by HDACs may be conserved in the development of different fibrotic diseases.

Conclusion. Counteracting the development of pancreatic fibrosis is a major and elusive therapeutic goal in the context of chronic pancreatitis. Our data revealed a potent antifibrotic effect of MS-275 treatment, which is mediated at least in part by suppression of TGFβ expression in acinar cells; however, it is possible that downregulation of additional factors contributes to the observed phenotype. In this context, it is worth mentioning that pancreatic expression of interleukins-1 and -6, interleukins known to promote autocrine and paracrine activation of PSC (Bynygeri et al., 2017), was reduced upon MS-275 treatment (Bombardo et al., 2017).

Collectively, our data suggest on one hand that class 1 HDAC activity is critical for the timely controlled epigenetic regulation of key signaling molecules driving the development of fibrosis in this organ. On the other hand, our data provide a new perspective on the cell types involved in regulating the process and highlights the possibility that acinar cells act as active mediators of pancreatic fibrosis.

These results harbor important implications to explore further the therapeutic potential of MS-275 in the context of chronic pancreatitis patients. Additional studies that include experimental models with increased severity of pancreatitis and autoimmune pancreatitis are warranted to define the effect of MS-275 in a broader spectrum of disease manifestations. Moreover, investigations using conditional knockout mouse models are needed to achieve a global understanding of the individual HDAC isoforms’ functions in the different cell types that are involved in the development of this disease.

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Authorship Contributions

Participated in research design: Bombardo, Graf, Sonda.

Conducted experiments: Bombardo, Chen, Malagola, Saponara.

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References


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SUPPLEMENTARY FIGURE LEGENDS

Supplemental figure 1. qPCR of inflammatory markers in pancreata at the indicated weeks of pancreatitis induction. F4/80, macrophage marker; Il6, interleukin 6; Mcp1, monocyte chemotactic protein 1. Results are average ± SD (n=5), *P < 0.05.

Supplemental figure 2. qPCR of HDAC isoforms in the pancreas upon 4 week treatment with cerulein in the presence or absence of MS-275. Data are expressed as ΔCt. Results are average ± SD (n=5).

Supplemental figure 3. qPCR of TGFβ1 and 2 expression in AR42J cells (A) and isolated acini (B) upon in vitro treatment with cerulein in the presence or absence of MS-275. Data are expressed as ΔCt. Results are average ± SD (n=5-6), *P < 0.05.
Figure S2
Figure S3