Vitamin A Inhibits Pancreatic Stellate Cell Activation: Implications for Treatment of Pancreatic Fibrosis

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Abbreviations:
ADH Alcohol dehydrogenase; ATRA All-trans retinoic acid; ANOVA Analysis of variance; ECL Enhanced chemiluminescence; ERK1/2 Extracellular-regulated kinases 1 and 2; FBS Fetal bovine serum; HSCs Hepatic stellate cells; IMDM Iscove’s Modified Dulbecco’s medium; JNK c-Jun N-terminal kinase; MAPK Mitogen-activated protein kinase; MAPKK MAPK kinase; MAPKKK MAPKK kinase; MKP-1 Mitogen-activated phosphatase-1; PSCs Pancreatic stellate cells; RAR Retinoic acid receptor; RXR Retinoid X receptor; ROLDH Retinol dehydrogenase; ROL retinol; 9-RA 9-cis retinoic acid; α-SMA Alpha-smooth muscle actin; SDS Sodium dodecyl sulfate; SV Sodium orthovanadate; TBS Tris buffered saline; TTBS Tris buffered saline and tween-20.

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**ABSTRACT:**

**Background & Aims:** Activated pancreatic stellate cells (PSCs) are implicated in the production of alcohol-induced pancreatic fibrosis. PSC activation is invariably associated with a loss of the cytoplasmic vitamin A (retinol) stores. Furthermore, retinol and ethanol are known to be metabolised by similar pathways. Our group and others have demonstrated that ethanol-induced PSC activation is mediated by the mitogen-activated protein kinase (MAPK) pathway, but the specific role of retinol and its metabolites all-trans retinoic acid (ATRA) and 9-cis retinoic acid (9-RA) in PSC quiescence/activation, or its influence on ethanol-induced PSC activation is not known. Therefore, the aims of this study were to i) examine the effects of retinol, ATRA and 9-RA on PSC activation; ii) determine whether retinol, ATRA and 9-RA influence MAPK signalling in PSCs; and iii) assess the effect of retinol supplementation on PSCs activated by ethanol. **Methods:** Cultured rat PSCs were incubated with retinol, ATRA or 9-RA for varying time periods and assessed for: i) proliferation; ii) expression of α-smooth muscle actin (α-SMA), collagen I, fibronectin and laminin; and iii) activation of MAPKs (ERK1/2, p38 kinase and JNK). The effect of retinol on PSCs treated with ethanol was also examined by incubating cells with ethanol in the presence or absence of retinol for 5 days, followed by assessment of α-SMA, collagen I, fibronectin and laminin expression. **Results:** Retinol, ATRA and 9-RA significantly inhibited: (i) cell proliferation, (ii) expression of α-SMA, collagen I, fibronectin and laminin, and iii) activation of all 3 classes of MAPKs. Furthermore, retinol prevented ethanol-induced PSC activation as indicated by inhibition of the ethanol-induced increase in α-SMA, collagen I, fibronectin and laminin expression. **Conclusions:** Retinol and its metabolites ATRA and 9-RA induce quiescence in culture-activated PSCs associated with a significant decrease in the activation of all 3 classes of MAPKs in PSCs. Ethanol-induced PSC activation is prevented by retinol supplementation.
INTRODUCTION:
Pancreatic fibrosis is a common histopathological feature of alcoholic chronic pancreatitis. Fibrogenesis involves a specific cell type in the pancreas, namely the pancreatic stellate cell (PSC). Recent studies have demonstrated that PSCs become activated [as indicated by proliferation, increased expression of the cytoskeletal protein alpha-smooth muscle actin (α-SMA) and synthesis of extracellular matrix proteins] when cultured on plastic or when exposed to ethanol and its oxidative metabolite acetaldehyde or to factors known to be upregulated during pancreatic injury such as proinflammatory cytokines, growth factors and oxidant stress.

PSCs in their quiescent (non-activated) state, store retinol as lipid droplets in their cytoplasm. Activation of PSCs is invariably associated with a loss of the retinol-containing droplets from their cytoplasm. Studies in other cell systems have established that retinol is a compound essential to normal cell biology. In particular, the metabolites of retinol, all-trans retinoic acid (ATRA) and 9-cis retinoic acid (9-RA) have been shown to mediate a number of cellular functions including proliferation, differentiation and protein synthesis. The metabolism of retinol has been best studied in the liver. It has been demonstrated in hepatic stellate cells (HSCs) that retinol enters the cell bound to cellular retinol binding protein (CRBP). It is then either esterified to retinyl ester for storage, or converted to retinaldehyde by retinol dehydrogenase (ROLDH) and subsequently to retinoic acid by retinaldehyde dehydrogenase (RALDH). Two forms of retinoic acid, ATRA and 9-RA serve as ligands for two families of nuclear receptors: retinoic acid receptors (RARα, β and γ) and retinoid X receptors (RXRα, β and γ). The RARs bind to ATRA with high affinity, whereas 9-RA is a bifunctional ligand which can bind to and activate both RARs and RXRs. Following ligand binding, these compounds interact with cis-acting DNA sequences called retinoic acid responsive elements (RAREs) in the promoter regions of target genes, thereby regulating gene expression. It is of interest that studies with HSCs have demonstrated that upon exposure to activating factors, the levels of retinoic acid and the expression of their receptors is decreased. Moreover, maintenance of the quiescent phenotype of HSCs has been shown to be dependent on adequate levels of retinoic acid and its receptors in the cells. A recent study by Jaster et al has reported decreased proliferation and collagen synthesis in PSCs upon exposure to ATRA. However, there has been no comprehensive study in the literature to date examining the effects of retinol and both its metabolites (ATRA and 9-RA) on PSC function.

A number of observations suggest a link between the metabolism of ethanol and retinol. Ethanol and its oxidative metabolite acetaldehyde are key activating factors of PSCs. The oxidation of ethanol to acetaldehyde is mediated by the enzyme alcohol dehydrogenase (ADH). Acetaldehyde is then further oxidised to acetic acid via aldehyde dehydrogenase (ALDH). It has been established that the retinol metabolising enzymes ROLDH and RALDH belong to the same family of alcohol and acetaldehyde-metabolising dehydrogenases respectively. Interestingly, studies have shown that rats chronically exposed to ethanol display significantly reduced retinoic acid levels in the liver. This is thought to be due to competitive inhibition by ethanol of retinol metabolism by ROLDH and ADH, since these enzymes can utilise both ethanol and retinol as substrates. Thus, retinoic acid depletion may be an important mechanism by which ethanol promotes stellate cell activation.

In recent studies, we and others have identified the mitogen-activated protein kinase (MAPK) pathway as a major cell signalling pathway mediating ethanol-induced and growth factor-induced PSC activation. MAPKs (which include extracellular-regulated kinases 1 and 2 [ERK1/2], N-terminal c-Jun kinase [JNK] and p38 kinase) play a major role in
regulating protein synthesis in mammalian cells. MAPKs modulate the activity of a number of downstream transcription factors and protein kinases by phosphorylation, thereby controlling gene expression and cell behaviour. Activation of MAPKs is a reversible process; they are inactivated by a group of protein phosphatases known as MAP kinase phosphatases (MKPs). These are dual-specificity phosphatases which have been shown to exert their effect by dephosphorylating tyrosine and threonine residues on MAPKs thereby leading to their inactivation. MAPK phosphatase-1 (MKP-1), a member of the MKP family, has recently been demonstrated to inactivate all 3 classes of MAPKs. It is interesting to note that recent studies have shown that MKP-1 expression is increased upon exposure to retinoic acid thereby resulting in a decrease in MAPK activity. However, whether retinol influences the MAPK signalling pathway or MKP-1 expression in stellate cells (whether from the pancreas or liver) is not known.

Therefore, the aims of this study were to: i) examine the effect of retinol and its metabolites ATRA and 9-RA on PSC activation, ii) determine whether retinol influences MAPK signalling and MKP-1 expression in PSCs, and iii) assess the influence of retinol on ethanol-treated PSCs.

METHODS:
Cell culture reagents, protease type XIV, all-trans retinol, all-trans retinoic acid, 9-cis retinoic acid were purchased from the Sigma Chemical Company (St Louis, MO, USA). Primers for ROLDH II, RARα, RXRα and RXRβ were obtained from Sigma Genosys Australia Pty Ltd. Sources of antibodies were as follows: α-SMA, fibronectin and laminin – Sigma; phospho-ERK1/2 and total ERK1/2, total p38 kinase and total JNK - Cell Signaling Technology (Beverly, MA, USA); phospho-p38 kinase and phospho-JNK - Promega (Sydney, Australia); MKP-1, RARα, RXRα and RXRβ - Santa Cruz Biotechnology (Santa Cruz, CA, USA); collagen type I - Rockland Immunochemicals (Gilbertsville, PA, USA); secondary goat anti-rabbit antibody - DAKO (Sydney, Australia).

Isolation and culture of PSCs
Rat pancreatic stellate cells were isolated by density gradient centrifugation as detailed previously. This technique results in a pure preparation of PSCs as evidenced by positive staining for stellate cell selective markers and negative staining for possible contaminants such as endothelial cells and macrophages. The yield of PSCs is 1.5 – 2 million cells per rat pancreas. Cells were cultured in Iscove’s Modified Dulbecco’s medium (IMDM) supplemented with 10% fetal bovine serum (FBS) at 37°C in humidified 95% air/5% CO2 atmosphere. All experiments were performed with culture-activated cells (passage 1-3) with the exception of experiments to assess the effect of retinol on freshly isolated (quiescent) cells.

Expression of retinol dehydrogenase and RAR and RXR receptor mRNA using RT-PCR
Expression of mRNA for retinol dehydrogenase II (ROLDH II) and the retinoic acid receptors RARα, RXRα and RXRβ was examined using reverse transcriptase-polymerase chain reaction (RT-PCR). Primer sequences were as follows:

Rat ROLDH II
Forward primer: 5’ ACC TGG CAT CTT ATC TGA AA 3’
Reverse primer: 5’ AGT CGA GTC AGC CTT GAG TA 3’

Rat RARα
Forward primer: 5’ CCC AGC CAC CAT TGA GAC 3’
Reverse primer: 5’ TAC ACC ATG TTC TTC TGG ATG C 3’

Rat RXRα
Forward primer: 5′ ATG AAG CGG GAA AGC TGT G 3′
Reverse primer: 5′ CAT GTT TGC CTC CAC GTA TG 3′

Rat RXRβ
Forward primer: 5′ TCA ACT CCA CAG TGT CGC TC 3′
Reverse primer: 5′ TAA ACC CCA TAG TGC TTG CC 3′

Treatment with Test Factors
1. Treatment of PSCs with retinol and retinoic acid
Culture-activated PSCs (passage 1-3) were exposed to retinol (10µM), ATRA (10µM) or 9-RA (10µM) for varying periods of time. Medium was changed every 48 hours. All manipulations were performed in subdued light. Cells incubated with culture medium with an equivalent amount of vehicle - DMSO 0.1% (for ATRA and 9-RA experiments) or DMSO 0.025% + 8mM ethanol (for retinol experiments) served as controls. Cell viability was assessed by trypan blue exclusion studies.

2. Treatment of PSCs with ethanol and retinol
To determine the effect of retinol on PSCs treated with ethanol, cells were incubated with 10µM retinol in the presence or absence of 50mM ethanol for 5 days. Medium was changed every 24 hours. At the end of the incubation period cell lysates were collected and expression of α-SMA, collagen I, fibronectin and laminin was assessed by western blotting. Cell viability was assessed as described earlier.

Assessment of PSC Function
1. Cell Proliferation
PSC proliferation was assessed by measuring the incorporation of [3H] thymidine into cellular DNA as previously described. In addition, cell counts were performed using a haemocytometer. Results are expressed as a percentage of control.

2. α-Smooth muscle actin and extracellular matrix protein expression
Levels of α-SMA, collagen I, fibronectin and laminin in PSCs were determined by western blotting of cell lysates as detailed previously, using a monoclonal mouse anti-α-SMA (1:200),3 or polyclonal rabbit anti-collagen I (1:1000), anti-fibronectin (1:1000) and anti-laminin (1:1000) antibodies.29, 30 A goat anti-mouse antibody (1:2000) or a goat anti-rabbit antibody (1:2000) was used as the secondary antibody. Expression of α-SMA, collagen I, fibronectin and laminin was detected by the enhanced chemiluminescence (ECL) technique using the Amersham enhanced ECL kit and quantified using densitometry (BioRad GelDoc Image One software).

3. Detection of MAPK activation
ERK1/2, p38 kinase and JNK activation in culture-activated PSCs exposed to retinol (10µM), ATRA (10µM) or 9-RA (10µM) for 4, 24 and 48 hours was assessed by western blotting of cell lysate proteins, as described previously.17, 28 To confirm that total MAPK levels were unchanged by the treatments, aliquots of the above cell lysates were subjected to western blotting analysis using rabbit polyclonal antibodies against ERK1/2, p38 kinase or JNK (1:1000), which recognise both the phosphorylated and non-phosphorylated forms of the enzymes; the blots were then analysed by densitometry.

Assessment of RAR and RXR Protein Expression in PSCs
RAR and RXR expression in PSCs treated with 10µM ATRA or 10µM 9-RA for 4, 24 and 48 hours was assessed by western blotting using polyclonal rabbit antibodies against RARα, RXRα or RXRβ (1:200).

Detection of MKP-1 Expression
MKP-1 expression in PSCs incubated with retinol, ATRA or 9-RA for 3, 6, and 24 hours was assessed by western blotting using a polyclonal rabbit anti-MKP-1 antibody (1:1000). To determine the role of MKP-1 in regulating the action of ATRA on MAPKs, we examined whether any observed inhibition of MAPKs by ATRA could be prevented by the protein phosphatase inhibitor sodium orthovanadate, a potent inhibitor of protein tyrosine phosphatases, which include MKP-1. The influence of this inhibitor on PSC function (cell proliferation and ECM protein expression) was also assessed. PSCs were treated with 10µM ATRA in the presence or absence of 1µM sodium orthovanadate (SV) for 48 hours. At the end of the incubation period, cell counts were performed using a haemocytometer and cell lysates were collected for assessment of MAPK activation and ECM protein expression by western blotting as described above.

Treatment of freshly isolated PSCs with retinol
To determine whether retinol could prevent/retard the transition of freshly isolated (quiescent) PSCs to their activated phenotype, freshly isolated cells were incubated in (IMDM) supplemented with 10% fetal bovine serum in the presence or absence of 10µM of retinol for 5 days. Medium was changed every 24 hours. At the end of the incubation period cell lysates were collected and expression of α-SMA and fibronectin was assessed as described above.

STATISTICS
Results are expressed as mean ± standard error; n represents the number of individual PSC preparations. Data were analysed using repeated measures analysis of variance (ANOVA). Fisher’s protected least significant difference (PLSD) was used for comparison of individual groups provided the F-test was significant. Analyses were performed using the Statview II® statistical software package.

RESULTS
Expression of retinol dehydrogenase II and retinoic acid receptors in PSCs
Using RT-PCR, retinol dehydrogenase II (ROLDH II), RARα, RXRα and RXRβ expression was observed in PSCs (Fig 1A and 1B, n = 3 separate cell preparations). The PCR products were 398bp, 195bp, 165bp and 175bp respectively, in agreement with previous reports.  

Retinol and its metabolites ATRA and 9-RA induce PSC quiescence
A) Cell proliferation
Retinol, ATRA and 9-RA reduced the proliferation of culture-activated PSCs (n = 4 separate cell preparations) as assessed by thymidine incorporation into cellular DNA as well as by direct cell counts. There was a significant reduction after 72 hours of incubation with retinol and this effect was sustained over 96 hours (Fig 2). Proliferation was also significantly reduced in the presence of ATRA and 9-RA (Fig 2). Notably, this reduction was observed at an earlier time point (48 hours) than that produced by retinol and this effect was sustained over 96 hours. We confirmed that the observed reduction in PSC proliferation
was not due to cytotoxicity of retinol, ATRA or 9-RA as evaluated by phase contrast microscopy and trypan blue exclusion studies (results not shown).

B) \(\alpha\)-SMA expression
Quiescence of PSCs after incubation with retinol (n = 3 separate cell preparations) was also demonstrated by significantly reduced \(\alpha\)-SMA expression after 72 hours, an effect that was sustained over 96 hours (Fig 3). Similar results were obtained with ATRA and 9-RA (Fig 3). ATRA was also associated with reduced \(\alpha\)-SMA expression at the earlier time point of 48 hours.

C) Extracellular matrix protein expression
ECM protein expression was significantly reduced in PSCs exposed to retinol or its metabolites (n = 3 separate cell preparations; Fig 4A, B and C). After 48 hours of incubation, retinol significantly decreased collagen I, fibronectin and laminin expression and this effect was sustained over 72 hours. Similar results were obtained for ATRA and 9-RA.

Retinol and its metabolites ATRA and 9-RA influence MAPK activation in PSCs

A) ERK 1/2 activation
Incubation of PSCs with retinol (n = 4 separate cell preparations) caused a significant decrease in ERK1/2 activation at 4 hours; this decrease was sustained over 48 hours (Fig 5). Similarly, ERK1/2 activation was reduced in the presence of ATRA and 9-RA (Fig 5). Western blotting of control and treated cell lysates for total ERK demonstrated that total ERK expression was unchanged by retinol, ATRA and 9-RA treatment [Densitometry data expressed as % of control (mean ± SEM): At 4h - Rol 100.5±5.8, ATRA 99.6±4.6; 9-RA 93.7±3.6; At 24h - Rol 96.7±9.2, ATRA 92.0±8.9; 9-RA 102.7±5.1; At 48h - Rol 107.75±10.6, ATRA 111.5±6.5; 9-RA 105.0±7.2]. Thus, these compounds appear to have a specific effect on ERK1/2 phosphorylation.

B) p38 kinase activation
Retinol significantly decreased p38 kinase activation after 48 hours of incubation (n = 4 separate cell preparations; Fig 6). Both ATRA and 9-RA also significantly reduced p38 kinase activation. However, this reduction was observed at the earlier time points of 4 and 24 hours and was sustained over 48 hours (Fig 6). Total p38 kinase expression was unchanged by the treatments. [Densitometry data expressed as % of control (mean ± SEM): At 4h - Rol 102.8±3.9, ATRA 94.2±3.96; 9-RA 90.57±4.8; At 24h - Rol 93.07±2.8, ATRA 98.6±0.48; 9-RA 99.2±2.8; At 48h - Rol 90.55±4.86, ATRA 95.4±3.0; 9-RA 102.6±5.8].

C) JNK activation
In contrast to the effect on ERK and p38 kinase activation, incubation of PSCs with retinol or 9-RA had no effect on JNK 2 (p54) activation (n = 4 separate cell preparations; Fig 7). However, ATRA significantly decreased JNK 2 (p54) activation after 4 hours and this decrease was sustained over 48 hours (Fig 7). Total JNK expression was not affected by Rol, ATRA or 9-RA treatment. [Densitometry data expressed as % of control (mean ± SEM): At 4h - Rol 95.4±10.0, ATRA 89.3±8.7; 9-RA 103.7±6.9; At 24h - Rol 123.8±11.5, ATRA 89.7±11.8; 9-RA 100.6±9.9; At 48h - Rol 117.3±13.4, ATRA 96.5±2.8; 9-RA 98.1±3.3].

Effect of retinol, ATRA and 9-RA on MKP-1 expression
ATRA significantly increased MKP-1 expression in PSCs (n = 3 separate cell preparations) at 6 hours and this increase was sustained over 24 hours (Fig 8). In contrast, retinol and 9-RA had no effect on MKP-1 expression (results not shown).

**Effect of the protein phosphatase inhibitor sodium orthovanadate on MAPK expression, cell proliferation and ECM protein expression in ATRA-treated PSCs**

To confirm whether MKP-1 plays a role in mediating the effect of ATRA on MAPK activation in PSCs, cells were treated with the protein phosphatase inhibitor sodium orthovanadate (n = 3 separate cell preparations). As shown in Figure 9, the ATRA-induced decrease in activation of ERK1/2, p38 kinase and JNK 2 was completely prevented in the presence of sodium orthovanadate (SV), suggesting that MKP-1 may mediate the ATRA-induced inhibition of MAPK activation. SV treatment of PSCs also prevented the ATRA-induced decrease in i) PSC proliferation (data expressed as % of control mean ± SEM: Control 100, ATRA 44.74 ± 8.72*, SV 85.35 ± 16.5, ATRA + SV 98.03 ± 26.8; *p<0.01 ATRA vs control) and ii) collagen I, fibronectin and laminin expression (Fig 10), further supporting the concept that the effects of ATRA on PSC function may be mediated via the MAPK pathway. Trypan blue exclusion studies confirmed cell viability in the presence of SV (results not shown).

**Effect of ATRA and 9-RA on RAR and RXR protein expression**

After 4 hours of incubation, both ATRA and 9-RA significantly increased RARα protein expression in PSCs (n = 3 separate cell preparations) and this effect was sustained over 24 and 48 hours (Fig 11A). 9-RA also increased the expression of RXRα and RXRβ at 4, 24 and 48 hours of incubation (Fig 11B).

**Effect of retinol on ethanol-induced PSC activation**

**A) α-SMA expression**

In order to determine whether retinol supplementation could prevent ethanol-induced PSC activation, α-SMA expression was assessed (n = 3 separate cell preparations, Fig 12). As expected, ethanol alone significantly increased α-SMA expression confirming our previously published results.2, 17 Retinol alone significantly reduced α-SMA expression confirming our results described in Fig 3. Importantly, retinol also significantly reduced α-SMA expression in PSCs treated with ethanol. Of particular interest was the finding that in the presence of ethanol, retinol was unable to fully exert its inhibitory effect when compared to retinol alone (compare Rol with Rol + E50, Fig 12).

**B) Extracellular matrix protein expression**

As illustrated in Fig 13, retinol significantly inhibited the expression of all 3 ECM proteins in PSCs (n = 3 separate cell preparations) and also prevented the ethanol-induced increase of collagen I, fibronectin and laminin expression in the cells. Interestingly, as with α-SMA expression, retinol was unable to fully exert its inhibitory effect on collagen I and laminin expression in the presence of ethanol, when compared to retinol alone.

**Effect of retinol on freshly isolated PSCs**

Incubation of freshly isolated PSCs with retinol for 5 days significantly decreased the expression of both α-SMA and fibronectin in the cells when compared to controls (Fig 14 A and B).
DISCUSSION
This study provides novel data demonstrating that vitamin A (retinol) induces quiescence in both freshly isolated and culture activated PSCs. We have also shown that the metabolites of vitamin A (ATRA and 9-RA) inhibit the activation of culture-activated PSCs. Notably, we have found that the above effects are associated with a significant decrease in the activation of all 3 classes of MAPKs, suggesting that vitamin A-induced quiescence is mediated via the MAPK pathway. Another novel finding of this study (of relevance to alcohol-induced fibrosis) is the prevention of ethanol-induced PSC activation by retinol. To the best of our knowledge, this aspect has not been previously studied in pancreatic or hepatic stellate cells.

The present study has also provided evidence indicating that PSCs possess the ability to convert retinol into its active metabolites. We have shown that cultured PSCs express the enzyme retinol dehydrogenase II (ROLDH II), an enzyme which is essential for the conversion of cellular retinol to retinoic acid. Our findings also suggest the presence of a functional retinoic acid signalling pathway in PSCs. We have identified mRNA for RAR and RXRs in the cells and moreover, have found that the expression of RAR and RXR proteins is induced by their ligands ATRA and 9-RA. The observed induction of receptor protein expression by their ligands in PSCs concurs with a previous report in cardiomyocytes, showing that both RAR and RXR are transcriptionally activated by ATRA and 9-RA.

The signalling pathways responsible for regulating PSC activation have been the focus of much interest in recent years, since therapeutic targeting of relevant signalling molecules may enable inhibition of PSC activation and prevention of fibrogenesis. Studies by our group and others have demonstrated that all 3 classes (ERK1/2, JNK and p38 kinase) of the MAPK pathway play an important role in mediating PSC activation upon exposure to profibrogenic factors. The ERK1/2 pathway has been shown to mediate PSC proliferation by increasing the activity of the transcription factor AP-1. Recently, our group and others have shown that the p38 kinase pathway plays a major role in mediating α-SMA expression in culture-activated and ethanol treated PSCs. Masamune et al. reported that both p38 kinase and JNK play an important role in mediating ECM protein synthesis in PSCs. Given the above, it would be reasonable to speculate that inhibition of PSC activation by vitamin A may be mediated by the inhibition of the MAPK pathway. The results of our study support this concept. We have demonstrated for the first time that retinol, ATRA and 9-RA significantly decrease the activation of ERK1/2 and p38 kinase in culture-activated PSCs after 4 hours of incubation and this decrease was sustained over 48 hours. In addition, ATRA (but not retinol or 9-RA) significantly decreased JNK 2 (p54) activation at 4, 24 and 48 hours. Our results with ATRA concur with those reported in human bronchial cells and hepatocytes. Both ERK1/2 and JNK MAPKs are known to regulate the activity of the transcription factor AP-1, which in turn is known to be an essential factor for the regulation of cell proliferation. Upon activation, AP-1 binds to a DNA sequence motif in target genes, thereby regulating their expression. A number of studies in other cell types have shown that increased AP-1 activity is essential in regulating cell proliferation (see review). It is of interest to note that upon exposure to vitamin A, AP-1 activity is decreased in a number of different cell types, and this is thought to be one of the mechanisms whereby vitamin A inhibits cell proliferation.

The mechanisms of MAPK inhibition (inactivation) by vitamin A are not yet determined. While activation of MAPK depends on phosphorylation of its tyrosine and threonine residues by upstream kinases, recent studies indicate that inactivation of MAPK is secondary to dephosphorylation via MAP kinase phosphatases (MKPs). MKP-1 has been
shown to inactivate all 3 classes of MAPKs. Interestingly, studies in other cell types have reported that treatment with vitamin A increases the expression of MKP-1 which in turn results in a decrease in MAPK activity. The current study has demonstrated that ATRA (but not retinol or 9-RA) induces the expression of MKP-1 in PSCs. This induction is evident after 6 hours of incubation and is sustained over 24 hours. We have further demonstrated that the protein phosphatase inhibitor sodium orthovanadate (known to inhibit phosphatases such as MKP-1) prevents the inhibition of MAPK activation in PSCs treated with ATRA. These findings support the concept that an increase in MKP-1 activity may play a role in the observed decrease in MAPK activation in PSCs exposed to ATRA. The mechanisms responsible for the retinol- and 9-RA- induced decrease in MAPK activation are unclear (given that these compounds did not increase MKP-1 activity in PSCs). It is possible that these compounds induce the activity of other phosphatases of the MAPK phosphatase family, as has been reported by Palm-Leis et al in cardiomyocytes.

It is now generally accepted that ethanol-induced PSC activation plays a major role in alcoholic pancreatic fibrosis. In view of our findings that retinol induces quiescence of PSCs in culture, we examined the effect of retinol on ethanol-treated PSCs. Our results have shown that retinol prevents ethanol-induced PSC activation as evidenced by a significant decrease in α-SMA expression, and more importantly, a significant decrease in the expression of the ECM proteins collagen I, fibronectin and laminin which are major components of fibrous tissue. In the course of these studies, an interesting observation was made with respect to the effect of retinol on PSC activation in the presence and absence of ethanol. In the presence of ethanol, retinol was found to be unable to exert its full inhibitory effect on α-SMA, collagen I, and laminin expression when compared to retinol alone (see Fig 12 Rol vs Rol+E50 for α-SMA expression and Fig 13 Rol vs Rol+E50 for collagen I and laminin expression). One possible explanation for this could be the competitive inhibition of retinol metabolism by ethanol, since the enzymes ROLDH and ADH can use both retinol and ethanol as substrates. There is evidence for the presence of both these enzymes in PSCs. The current study has demonstrated the expression of mRNA for ROLDH in PSCs, while previous studies by our group have demonstrated alcohol dehydrogenase (ADH) enzyme activity (inducible after exposure to ethanol) in PSCs. A decrease in retinol metabolism in ethanol-treated PSCs would lead to lower retinoic acid levels in these cells when compared with PSCs treated with retinol alone.

In conclusion, this study is the first to demonstrate that vitamin A and/or its metabolites ATRA and 9-RA induce quiescence in both freshly isolated and culture-activated PSCs. Induction of PSC quiescence is associated with a decrease in the activation of all 3 classes of MAPKs. Furthermore, we have demonstrated that retinol supplementation prevents ethanol-induced activation of PSCs. These findings suggest that vitamin A may be a potentially useful antifibrotic agent in chronic pancreatitis via its ability to inhibit PSC activation and consequently reduce their synthesis of ECM proteins.

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Competing Interest: None declared
References


FIGURE LEGENDS

FIGURE 1A and B: Expression of ROLDH II, RAR and RXR in PSCs. A: Total RNA was extracted from 3 separate cell preparations and analysed for ROLDH II expression by RT-PCR. An RNA ladder was run in lane 1 (Std) to determine the size of the PCR products observed. Lane 2 contained rat liver cDNA as a positive control. Lane 3 contained a PCR product for ROLDH II from rat PSCs. B: RARα, RXRα and RXRβ expression was analysed in total RNA obtained from 3 separate cell preparations by RT-PCR. Lane 1 contains an RNA ladder (Std) to determine the size of the PCR products observed. Lanes 2, 3 and 4 contain PCR products for RARα, RXRα and RXRβ from rat PSCs.

FIGURE 2: Effect of retinol and its metabolites on PSC proliferation. (A) [3H] thymidine incorporation studies and (B) cell count data demonstrating that incubation of PSCs with retinol (Rol) decreased PSC proliferation significantly when compared to controls after 72 hours and that this effect was sustained over 96 hours. ATRA and 9-RA also significantly decreased PSC proliferation after 48 hours when compared to controls and this effect was sustained over 96 hours (n = 3 separate cell preparations; **p < 0.01, *p < 0.001).

FIGURE 3: Effect of retinol and its metabolites on α-SMA expression in PSCs. (A) Representative western blots showing α-SMA expression in PSCs incubated with culture medium alone (Cont), Rol, ATRA or 9-RA for 48, 72 and 96 hours. (B) Densitometry analysis of all western blots showing a significant decrease in α-SMA expression, at 72 and 96 hours in PSCs treated with Rol and at 48, 72 and 96 hours in PSCs treated with ATRA, compared to controls (n = 3 separate cell preparations; *p < 0.01, **p < 0.02, †p < 0.03).

FIGURE 4 A, B and C: Effect of retinol and its metabolites on extracellular matrix protein expression in culture-activated PSCs. Representative western blots and densitometry analysis showing a significant decrease in the expression of (A) collagen I, (B) fibronectin and (C) laminin, after treatment with Rol, ATRA or 9-RA for 48 and 72 hours when compared to controls (n = 3 separate cell preparations; *p < 0.005 for collagen I; *p < 0.001 for fibronectin; *p < 0.004, **p < 0.005, #p < 0.003, †p < 0.001 for laminin).

FIGURE 5: Effect of retinol and its metabolites on ERK1/2 activation in culture-activated PSCs. (A) Representative western blots showing ERK1/2 activation (p-ERK1/2) and total ERK1/2 expression in PSCs incubated with Rol, ATRA or 9-RA for 4, 24 and 48 hours. B: Densitometry analysis of all western blots showing a significant decrease in ERK1/2 activation in PSCs treated with Rol, ATRA or 9-RA after 4, 24 and 48 hours compared to controls (n = 4 separate cell preparations; (*p < 0.001, **p < 0.02, # p < 0.04). Rol, ATRA or 9-RA treatment had no significant effect on total ERK 1/2 levels in PSCs.

FIGURE 6: Effect of retinol and its metabolites on p38 kinase activation in culture-activated PSCs. A: Representative western blots showing p38 kinase activation in PSCs incubated with Rol, ATRA or 9-RA for 4, 24 and 48 hours. B: Densitometry analysis of all western blots showing a significant decrease in p38 kinase activation in PSCs treated with Rol after 48 hours compared to controls. Treatment with ATRA or 9-RA also resulted in a significant decrease in p38 kinase activation after 4 hours and was sustained after 48 hours compared to controls (n = 4 separate cell preparations; *p < 0.01, **p < 0.02, *p < 0.03). Rol, ATRA or 9-RA treatment had no significant effect on total p38 kinase levels in PSCs.
FIGURE 7: Effect of retinol and its metabolites on JNK activation in culture-activated PSCs. (A) Representative western blots showing JNK activation in PSCs incubated with Rol, ATRA or 9-RA for 4, 24 and 48 hours. (B) Densitometry analysis of all western blots showing a significant decrease in JNK 2 (p54) activation after 4, 24 and 48 hours treatment with ATRA compared to controls (n = 3 separate cell preparations; *p < 0.01, **p < 0.005). Both Rol and 9-RA had no effect on JNK 2 (p54) activation. Total JNK 2 levels were unchanged by Rol, ATRA or 9-RA treatment.

FIGURE 8: Effect of ATRA on MKP-1 expression in culture-activated PSCs. (A) Representative western blot showing MKP-1 expression in PSCs incubated with culture medium alone (Cont) or ATRA for 3, 6 and 24 hours. (B) Densitometry analysis of western blots showing a significant increase in MKP-1 expression after 6 hours of treatment with ATRA; and this effect was sustained over 24 hours compared to controls (n = 3 separate cell preparations; *p < 0.005). Both Rol and 9-RA had no effect on MKP-1 expression (data not shown).

FIGURE 9: Effect of sodium orthovanadate (SV) on MAPK activation in ATRA treated cells. Representative western blots and densitometry analysis showing a significant decrease in activation of all 3 classes of MAPK in PSCs treated with ATRA for 24h. This decrease was prevented in the presence of sodium orthovanadate (*p < 0.001, **p < 0.02 ATRA vs control; #p < 0.003, ##p < 0.02 ATRA vs ATRA+SV; n = 3 separate cell preparations). Total MAPK levels were unchanged by the treatments.

FIGURE 10: Effect of sodium orthovanadate on ATRA-induced inhibition of ECM protein expression in PSCs. Representative western blots and densitometry analysis showing a significant decrease in collagen I, fibronectin and laminin expression in PSCs treated with ATRA for 48h and prevention of this decrease in the presence of sodium orthovanadate (*p < 0.02, **p < 0.03 ATRA vs control; #p < 0.02, ##p < 0.04 ATRA vs ATRA+SV; n = 3 separate cell preparations).

FIGURE 11A and B: Effect of ATRA and 9-RA on RAR and RXR receptor expression in culture-activated PSCs. (A) Representative western blots and densitometry analysis showing a significant increase in RARα protein expression in PSCs treated with ATRA for 4, 24 and 48 hours (*p < 0.04, **p < 0.03, ***p < 0.02; n = 3 separate cell preparations) (B) Representative western blots and densitometry analysis showing a significant increase in RXRα and RXRβ protein expression in PSCs treated with 9-RA for 4, 24 and 48 hours (***p < 0.03, ***p < 0.02, #p < 0.003, ♦p < 0.004, •p < 0.005; n = 3 separate cell preparations). Note that both RXR α and RXR β were represented by a number of protein bands with increased intensity suggesting the presence of multiple activated isoforms.

FIGURE 12: Effect of retinol on ethanol-induced α-SMA expression in PSCs. (A) Representative western blot showing α-SMA expression in PSCs incubated with culture medium alone (Cont) or with 50mM ethanol (E50) in the presence or absence of retinol for 5 days. B: Densitometry analysis of western blots showing a significant increase in α-SMA expression in PSCs when treated with ethanol (n = 3 separate cell preparations; *p < 0.01) when compared to controls. Retinol significantly decreased basal α-SMA expression (n = 3 separate cell preparations; **p < 0.02 Rol vs Cont) and prevented the ethanol-induced increase in α-SMA expression in PSCs (n =3 separate cell preparations; #p < 0.01 E50 vs E50 + Rol).

FIGURE 13: Effect of retinol on ethanol-induced extracellular matrix protein synthesis in PSCs. (A) Representative western blot showing collagen I, fibronectin and
laminin expression in PSCs incubated with culture medium alone (Cont) or with 50mM ethanol (E50) in the presence or absence of retinol for 5 days. (B) Densitometry analysis of western blots showing a significant increase in collagen I, fibronectin and laminin expression in PSCs treated with ethanol when compared to controls (n = 3 separate cell preparations; *p < 0.005 E50 vs Cont). Retinol significantly decreased basal collagen I, fibronectin and laminin levels (n = 3 separate cell preparations; #p < 0.05 Rol vs Cont) and prevented the ethanol-induced increase in ECM protein expression (n = 3 separate cell preparations; †p < 0.05 E50 vs E50 + Rol).

FIGURE 14: Effect of retinol on freshly isolated PSC activation. Representative western blots and densitometry analysis showing a significant decrease in the expression of (A) α-SMA and (B) fibronectin, in freshly isolated PSCs treated with retinol for 5 days when compared to controls (n = 4 separate cell preparations; *p < 0.001).
FIGURE 1:

A) 

Lane:  DNA Std (bp) 21 3 
RolDH II (398bp) 
Liver +ve cont 

B) 

Lane:  DNA Std (bp) 1 2 3 4 
RAR\(\alpha\) (195bp) RXR\(\alpha\) (165bp) RXR\(\beta\) (175bp) 
PSCs
FIGURE 2:

A) 

$^3$H-Thymidine Incorporation into DNA (% of Control)

B) 

Cell Counts (% of Control)
FIGURE 3:

A)  

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B)  

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<th>Rol</th>
<th>Cont</th>
<th>ATRA</th>
<th>Cont</th>
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FIGURE 4A:

Collagen I 150KDa

48hrs

72 hrs

Cont Rol Cont ATRA Cont 9-RA

Densitometry Units (% of Control)

0 20 40 60 80 100 120

48h 72h

* * *
FIGURE 4B:

48 hrs
Fibronectin → 250KDa

72 hrs
Fibronectin → 250KDa
Cont  Rol Cont ATRA Cont 9-RA

Densitometry Units (% of Control)

48h 72h
FIGURE 4C:

48 hrs
Laminin

250KDa

72 hrs
Laminin

250KDa

Cont  Rol  Cont  ATRA  Cont  9-RA

Densitometry Units (% of Control)

Cont  Rol  ATRA  9-RA

48h  72h
FIGURE 5:

A) Rol

p-ERK1/2 →
Total-ERK1/2 →
Cont 4h 24h 48h Cont 24h 48h

ATRA

p-ERK1/2 →
Total-ERK1/2 →
Cont 4h 24h 48h

9-RA

p-ERK1/2 →
Total-ERK1/2 →
Cont 4h 24h 48h

B) Densitometry Units (% of Control)

Cont Rol ATRA 9-RA

4h 24h 48h
FIGURE 6:

A)

Rol

p-p38 kinase

Total-p38 kinase

Cont 4h 24h 48h

ATRA

p-p38 kinase

Total-p38 kinase

Cont 4h 24h 48h

9-RA

p-p38 kinase

Total-p38 kinase

Cont 4h 24h 48h

B)

Densitometry Units (% of Control)

Cont Rol ATRA 9-RA

4h 24h 48h
FIGURE 7:

A)

- Rol
  - p-JNK 2 (p54)
  - Total-JNK 2 (p54)
  - Cont, 4h, 24h, 48h
- ATRA
  - p-JNK 2 (p54)
  - Total-JNK 2 (p54)
  - Cont, 4h, 24h, 48h
- 9-RA
  - p-JNK 2 (p54)
  - Total-JNK 2 (p54)
  - Cont, 4h, 24h, 48h

B)

- Densitometry Units (% of Control)
  - Cont, Rol, ATRA, 9-RA
  - 4h, 24h, 48h
FIGURE 8:

A) ATRA

MKP-1

Cont 3h 6h 24h

Densitometry Units (% of Control)

B)
FIGURE 9:

- **p-ERK1/2**
- **Total-ERK1/2**
- **p-p38 kinase**
- **Total-p38 kinase**
- **p-JNK 2 (p54)**
- **Total-JNK 2 (p54)**

**Graph:**
- Densitometry Units (% of Control)

- **Cont**
- **ATRA**
- **SV**
- **ATRA + SV**

Legend:
- ERK1/2
- p38 kinase
- JNK
FIGURE 10:

Collagen I

Fibronectin

Laminin

Cont ATRA SV ATRA+SV

Densitometry Units (% of Control)

Cont ATRA SV ATRA+SV

Collagen I Fibronectin Laminin

** * # #

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FIGURE 11A:

A)
FIGURE 11B:

RXRα

Cont 9-RA 9-RA 9-RA
4h 24h 48h

RXRβ

Cont 9-RA 9-RA 9-RA
4h 24h 48h

Densitometry Units (% of Control)

Cont 4h 24h 48h

0 50 100 150 200 250

** *** #

RXRα RXRβ
FIGURE 12:

A) α-SMA

B) Densitometry Units (% of Control)

Cont  E50  Rol  Rol + E50

Cont  E50  Rol  E50 + Rol
FIGURE 13:

A) Collagen I 150KDa

Fibronectin 250KDa

Laminin 250KDa

B) Densitometry Units (% of Control)
FIGURE 14:

A) $\alpha$-SMA

B) Fibronectin

Densitometry Units (% of Control)